



## NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAME

### BACKGROUND OF THE INVENTION

The invention generally relates to nucleic acids and polypeptides. More particularly, the invention relates to nucleic acids encoding novel G-protein coupled receptor (GPCR) polypeptides, as well as vectors, host cells, antibodies, and recombinant methods for producing these nucleic acids and polypeptides.

### SUMMARY OF THE INVENTION

The invention is based in part upon the discovery of nucleic acid sequences encoding novel polypeptides. These nucleic acids and polypeptides, as well as derivatives, homologs, analogs and fragments thereof, will hereinafter be collectively designated as "GPCRX" nucleic acid or polypeptide sequences.

In one aspect, the invention provides an isolated GPCRX nucleic acid molecule encoding a GPCRX polypeptide that includes a nucleic acid sequence that has identity to the nucleic acids disclosed in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39. In some embodiments, the GPCRX nucleic acid molecule will hybridize under stringent conditions to a nucleic acid sequence complementary to a nucleic acid molecule that includes a protein-coding sequence of a GPCRX nucleic acid sequence. The invention also includes an isolated nucleic acid that encodes a GPCRX polypeptide, or a fragment, homolog, analog or derivative thereof. For example, the nucleic acid can encode a polypeptide at least 80% identical to a polypeptide comprising the amino acid sequences of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40. The nucleic acid can be, for example, a genomic DNA fragment or a cDNA molecule that includes the nucleic acid sequence of any of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39. Also included in the invention is an oligonucleotide, e.g., an oligonucleotide which includes at least 6 contiguous nucleotides of a GPCRX nucleic

acid (e.g., SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39) or a complement of said oligonucleotide.

Also included in the invention are substantially purified GPCR<sub>X</sub> polypeptides (e.g., SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40). In certain embodiments, the GPCR<sub>X</sub> polypeptides include an amino acid sequence that is substantially identical to the amino acid sequence of a human GPCR<sub>X</sub> polypeptide.

The invention also features antibodies that immunoselectively bind to GPCR<sub>X</sub> polypeptides, or fragments, homologs, analogs or derivatives thereof.

In another aspect, the invention includes pharmaceutical compositions that include therapeutically- or prophylactically-effective amounts of a therapeutic and a pharmaceutically-acceptable carrier. The therapeutic can be, e.g., a GPCR<sub>X</sub> nucleic acid, a GPCR<sub>X</sub> polypeptide, or an antibody specific for a GPCR<sub>X</sub> polypeptide. In a further aspect, the invention includes, in one or more containers, a therapeutically- or prophylactically-effective amount of this pharmaceutical composition.

In a further aspect, the invention includes a method of producing a polypeptide by culturing a cell that includes a GPCR<sub>X</sub> nucleic acid, under conditions allowing for expression of the GPCR<sub>X</sub> polypeptide encoded by the DNA. If desired, the GPCR<sub>X</sub> polypeptide can then be recovered.

In another aspect, the invention includes a method of detecting the presence of a GPCR<sub>X</sub> polypeptide in a sample. In the method, a sample is contacted with a compound that selectively binds to the polypeptide under conditions allowing for formation of a complex between the polypeptide and the compound. The complex is detected, if present, thereby identifying the GPCR<sub>X</sub> polypeptide within the sample.

The invention also includes methods to identify specific cell or tissue types based on their expression of a GPCR<sub>X</sub>.

Also included in the invention is a method of detecting the presence of a GPCR<sub>X</sub> nucleic acid molecule in a sample by contacting the sample with a GPCR<sub>X</sub> nucleic acid probe or primer, and detecting whether the nucleic acid probe or primer bound to a GPCR<sub>X</sub> nucleic acid molecule in the sample.

In a further aspect, the invention provides a method for modulating the activity of a GPCR<sub>X</sub> polypeptide by contacting a cell sample that includes the GPCR<sub>X</sub> polypeptide with a compound that binds to the GPCR<sub>X</sub> polypeptide in an amount sufficient to modulate the activity of said polypeptide. The compound can be, e.g., a small molecule, such as a nucleic

acid, peptide, polypeptide, peptidomimetic, carbohydrate, lipid or other organic (carbon containing) or inorganic molecule, as further described herein.

Also within the scope of the invention is the use of a therapeutic in the manufacture of a medicament for treating or preventing disorders or syndromes including, *e.g.*,

5 developmental diseases; MHCII and III diseases (immune diseases); taste and scent detectability disorders; Burkitt's lymphoma; corticoneurogenic disease; signal transduction pathway disorders; metabolic pathway disorders; retinal diseases including those involving photoreception; cell growth rate disorders; cell shape disorders; metabolic disorders; feeding disorders; control of feeding; the metabolic syndrome X; wasting disorders associated with

10 chronic diseases; obesity; potential obesity due to over-eating or metabolic disturbances; potential disorders due to starvation (lack of appetite); diabetes; noninsulin-dependent diabetes mellitus (NIDDM1); infectious disease (bacterial, fungal, protozoal and viral infections), particularly infections caused by HIV-1 or HIV-2); pain; cancer (including but not limited to neoplasm adenocarcinoma, lymphoma, prostate cancer, and uterus cancer);

15 cancer-associated cachexia; anorexia; bulimia; asthma; Parkinson's disease; acute heart failure; hypotension; hypertension; urinary retention; osteoporosis; Crohn's disease; multiple sclerosis; Albright Hereditary Osteodystrophy; angina pectoris; myocardial infarction; ulcers; allergies; benign prostatic hypertrophy; and psychotic and neurological disorders including anxiety, schizophrenia, manic depression, delirium, dementia, neurodegenerative

20 disorders; Alzheimer's disease; severe mental retardation; Dentatorubro-pallidoluysian atrophy (DRPLA); Hypophosphatemic rickets; autosomal dominant (2) acrocallosal syndrome and dyskinesias, such as Huntington's disease or Gilles de la Tourette syndrome; immune disorders; adrenoleukodystrophy; congenital adrenal hyperplasia; hemophilia; hypercoagulation; idiopathic thrombocytopenic purpura; autoimmune disease;

25 immunodeficiencies; transplantation; Von Hippel-Lindau (VHL) syndrome; stroke; tuberous sclerosis; hypercalcaemia; cerebral palsy; epilepsy; Lesch-Nyhan syndrome; ataxia-telangiectasia; Leukodystrophies; behavioral disorders; addiction; neuroprotection; cirrhosis; transplantation; systemic lupus erythematosus; emphysema; scleroderma; ARDS; renal artery stenosis; interstitial nephritis; glomerulonephritis; polycystic kidney disease; renal tubular

30 acidosis; IgA nephropathy; cardiomyopathy; atherosclerosis; congenital heart defects; aortic stenosis; atrial septal defect (ASD); atrioventricular (A-V) canal defect; ductus arteriosus; pulmonary stenosis; subaortic stenosis; ventricular septal defect (VSD); valve diseases; scleroderma; fertility; pancreatitis; endocrine dysfunctions; growth and reproductive disorders; inflammatory bowel disease; diverticular disease; leukodystrophies; graft vesus

host; hyperthyroidism; endometriosis; hematopoietic disorders and/or other pathologies and disorders of the like. The therapeutic can be, *e.g.*, a GPCR<sub>X</sub> nucleic acid, a GPCR<sub>X</sub> polypeptide, or a GPCR<sub>X</sub>-specific antibody, or biologically-active derivatives or fragments thereof.

5           For example, the compositions of the present invention will have efficacy for treatment of patients suffering from the diseases and disorders listed above and/or other pathologies and disorders.

          The polypeptides can be used as immunogens to produce antibodies specific for the invention and as vaccines. They can also be used to screen for potential agonist and  
10       antagonist compounds. For example, a cDNA encoding GPCR<sub>X</sub> may be useful in gene therapy, and GPCR<sub>X</sub> may be useful when administered to a subject in need thereof. By way of nonlimiting example, the compositions of the present invention will have efficacy for treatment of patients suffering the diseases and disorders listed above and/or other pathologies and disorders.

15           The invention further includes a method for screening for a modulator of disorders or syndromes including, *e.g.*, diseases and disorders listed above and/or other pathologies and disorders and those disorders related to cell signal processing and metabolic pathway modulation. The method includes contacting a test compound with a GPCR<sub>X</sub> polypeptide and determining if the test compound binds to said GPCR<sub>X</sub> polypeptide. Binding of the test  
20       compound to the GPCR<sub>X</sub> polypeptide indicates the test compound is a modulator of activity, or of latency or predisposition to the aforementioned disorders or syndromes.

          Also within the scope of the invention is a method for screening for a modulator of activity, or of latency or predisposition to an disorders or syndromes including the diseases and disorders listed above and/or other pathologies and disorders or other disorders related to  
25       cell signal processing and metabolic pathway modulation by administering a test compound to a test animal at increased risk for the aforementioned disorders or syndromes. The test animal expresses a recombinant polypeptide encoded by a GPCR<sub>X</sub> nucleic acid. Expression or activity of GPCR<sub>X</sub> polypeptide is then measured in the test animal, as is expression or activity of the protein in a control animal which recombinantly-expresses GPCR<sub>X</sub>  
30       polypeptide and is not at increased risk for the disorder or syndrome. Next, the expression of GPCR<sub>X</sub> polypeptide in both the test animal and the control animal is compared. A change in the activity of GPCR<sub>X</sub> polypeptide in the test animal relative to the control animal indicates the test compound is a modulator of latency of the disorder or syndrome.



In yet another aspect, the invention includes a method for determining the presence of or predisposition to a disease associated with altered levels of a GPCR<sub>X</sub> polypeptide, a GPCR<sub>X</sub> nucleic acid, or both, in a subject (*e.g.*, a human subject). The method includes measuring the amount of the GPCR<sub>X</sub> polypeptide in a test sample from the subject and  
5 comparing the amount of the polypeptide in the test sample to the amount of the GPCR<sub>X</sub> polypeptide present in a control sample. An alteration in the level of the GPCR<sub>X</sub> polypeptide in the test sample as compared to the control sample indicates the presence of or predisposition to a disease in the subject. Preferably, the predisposition includes diseases and disorders listed above and/or other pathologies and disorders. Also, the expression levels of  
10 the new polypeptides of the invention can be used in a method to screen for various cancers as well as to determine the stage of cancers.

In a further aspect, the invention includes a method of treating or preventing a pathological condition associated with a disorder in a mammal by administering to the subject a GPCR<sub>X</sub> polypeptide, a GPCR<sub>X</sub> nucleic acid, or a GPCR<sub>X</sub>-specific antibody to a  
15 subject (*e.g.*, a human subject), in an amount sufficient to alleviate or prevent the pathological condition. In preferred embodiments, the disorder includes the diseases and disorders listed above and/or other pathologies and disorders.

In yet another aspect, the invention can be used in a method to identify the cellular receptors and downstream effectors of the invention by any one of a number of techniques  
20 commonly employed in the art. These include, but are not limited to, the two-hybrid system, affinity purification, co-precipitation with antibodies or other specific-interacting molecules.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can  
25 be used in the practice or testing of the present invention, suitable methods and materials are described below. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In the case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.  
30 Other features and advantages of the invention will be apparent from the following detailed description and claims.

## DETAILED DESCRIPTION OF THE INVENTION

The invention is based, in part, upon the discovery of novel nucleic acid sequences that encode novel polypeptides. The nucleic acids, and their encoded polypeptides, are collectively designated herein as "GPCRX".

5           The novel GPCRX nucleic acids of the invention include the nucleic acids whose sequences are provided in Table 1 (at the end of the Detailed Description), or a fragment, derivative, analog or homolog thereof. The novel GPCRX proteins of the invention include the protein fragments whose sequences are provided in Table 1. All of the sequences listed in Table 1 have a high degree of homology to known GPCR sequences. Exemplary homology  
10       for the sequences is provided in the provisional applications from which the present application claims priority. This homology data are incorporated herein by reference in their entirety. Within the scope of this invention is a method of using these nucleic acids and peptides in the treatment or prevention of a disorder related to cell signaling or metabolic pathway modulation.

15           G-Protein Coupled Receptor proteins ("GPCRs") have been identified as a large family of G protein-coupled receptors in a number of species. These receptors share a seven transmembrane domain structure with many neurotransmitter and hormone receptors, and are likely to underlie the recognition and G-protein-mediated transduction of various signals. Human GPCRs generally do not contain introns and belong to four different gene  
20       subfamilies, displaying great sequence variability. These genes are dominantly expressed in olfactory epithelium. See, *e.g.*, Ben-Arie et al., *Hum. Mol. Genet.* 1994 3:229-235; and, Online Mendelian Inheritance in Man ("OMIM") entry # 164342 (<http://www.ncbi.nlm.nih.gov/entrez/dispomim.cgi?>).

          The olfactory receptor ("OR") gene family constitutes one of the largest GPCR  
25       multigene families and is distributed among many chromosomal sites in the human genome. See Rouquier et al., *Hum. Mol. Genet.* 7:1337-45 (1998); Malnic et al., *Cell* 96:713-23 (1999). Olfactory receptors constitute the largest family among G protein-coupled receptors, with up to 1000 members expected. See Vanderhaeghen et al., *Genomics* 39:239-46 (1997); Xie et al., *Mamm. Genome* 11:1070-78 (2000); Issel-Tarver et al., *Proc. Natl. Acad. Sci. USA*  
30       93:10897-902 (1996). The recognition of odorants by olfactory receptors is the first stage in odor discrimination. See Krautwurst et al., *Cell* 95:917-26 (1998); Buck et al., *Cell* 65:175-87 (1991). Many ORs share some characteristic sequence motifs and have a central variable region corresponding to a putative ligand binding site. See Issel-Tarver et al., *Proc. Natl. Acad. Sci. USA* 93:10897-902 (1996).

Other examples of seven membrane spanning proteins that are related to GPCRs are chemoreceptors. See Thomas et al., *Gene* 178(1-2):1-5 (1996). Chemoreceptors have been identified in taste, olfactory, and male reproductive tissues. See *id.*; Walensky et al., *J. Biol. Chem.* 273(16):9378-87 (1998); Parmentier et al., *Nature* 355(6359):453-55 (1992); Asai et al., *Biochem. Biophys. Res. Commun.* 221(2):240-47 (1996).

The GPCR<sub>X</sub> nucleic acids of the invention encoding GPCR-like proteins include the nucleic acids whose sequences are provided herein, or fragments thereof. The invention also includes mutant or variant nucleic acids where any one or more bases may be changed from the corresponding base shown herein while still encoding a protein that maintains its GPCR-like activities and physiological functions, or a fragment of such a nucleic acid. The invention further includes nucleic acids whose sequences are complementary to those just described, including nucleic acid fragments that are complementary to any of the nucleic acids just described. The invention additionally includes nucleic acids or nucleic acid fragments, or complements thereto, whose structures include chemical modifications. Such modifications include, by way of nonlimiting example, modified bases, and nucleic acids whose sugar phosphate backbones are modified or derivatized. These modifications are carried out at least in part to enhance the chemical stability of the modified nucleic acid, such that they may be used, for example, as antisense binding nucleic acids in therapeutic applications in a subject.

The GPCR<sub>X</sub> proteins of the invention include the GPCR-like proteins whose sequences are provided herein. The invention also includes mutant or variant proteins any of whose residues may be changed from the corresponding residue shown herein while still encoding a protein that maintains its GPCR-like activities and physiological functions, or a functional fragment thereof. The invention further encompasses antibodies and antibody fragments, such as F<sub>ab</sub> or (F<sub>ab</sub>)<sub>2</sub>, that bind immunospecifically to any of the proteins of the invention.

The GPCR<sub>X</sub> nucleic acids and proteins are useful in potential therapeutic applications implicated in various GPCR-related pathological disorders and/or OR-related pathological disorders, described further below. For example, a cDNA encoding the GPCR (or olfactory-receptor) like protein may be useful in gene therapy, and the receptor-like protein may be useful when administered to a subject in need thereof. The nucleic acids and proteins of the invention are also useful in potential therapeutic applications used in the treatment of developmental diseases; MHCII and III diseases (immune diseases); taste and scent detectability disorders; Burkitt's lymphoma; corticoneurogenic disease; signal transduction

pathway disorders; metabolic pathway disorders; retinal diseases including those involving  
 photoreception; cell growth rate disorders; cell shape disorders; metabolic disorders; feeding  
 disorders; control of feeding; the metabolic syndrome X; wasting disorders associated with  
 chronic diseases; obesity; potential obesity due to over-eating or metabolic disturbances;  
 5 potential disorders due to starvation (lack of appetite); diabetes; noninsulin-dependent  
 diabetes mellitus (NIDDM1); infectious disease; bacterial, fungal, protozoal and viral  
 infections (particularly infections caused by HIV-1 or HIV-2); pain; cancer (including but not  
 limited to neoplasm, adenocarcinoma, lymphoma, prostate cancer, and uterus cancer); cancer-  
 associated cachexia; anorexia; bulimia; asthma; Parkinson's disease; acute heart failure;  
 10 hypotension; hypertension; urinary retention; osteoporosis; Crohn's disease; multiple  
 sclerosis; Albright hereditary osteodystrophy; angina pectoris; myocardial infarction; ulcers;  
 allergies; benign prostatic hypertrophy; psychotic and neurological disorders (including  
 anxiety, schizophrenia, manic depression, delirium, and dementia); neurodegenerative  
 disorders; Alzheimer's disease; severe mental retardation; dentatorubro-pallidoluysian  
 15 atrophy (DRPLA); hypophosphatemic rickets; autosomal dominant (2) acrocallosal syndrome  
 and dyskinesias, such as Huntington's disease or Gilles de la Tourette syndrome; immune  
 disorders; adrenoleukodystrophy; congenital adrenal hyperplasia; hemophilia;  
 hypercoagulation; idiopathic thrombocytopenic purpura; autoimmune disease;  
 immunodeficiencies; transplantation; Von Hippel-Lindau (VHL) syndrome; stroke; tuberous  
 20 sclerosis; hypercalcaemia; cerebral palsy; epilepsy; Lesch-Nyhan syndrome; ataxia-  
 telangiectasia; leukodystrophies; behavioral disorders; addiction; neuroprotection; cirrhosis;  
 transplantation; systemic lupus erythematosus; emphysema; scleroderma; ARDS; renal artery  
 stenosis; interstitial nephritis; glomerulonephritis; polycystic kidney disease; renal tubular  
 acidosis; IgA nephropathy; cardiomyopathy; atherosclerosis; congenital heart defects; aortic  
 25 stenosis ; atrial septal defect (ASD); atrioventricular (A-V) canal defect; ductus arteriosus;  
 pulmonary stenosis; subaortic stenosis; ventricular septal defect (VSD); valve diseases;  
 scleroderma; fertility; pancreatitis; endocrine dysfunctions; growth and reproductive  
 disorders; inflammatory bowel disease; diverticular disease; leukodystrophies; graft versus  
 host; hyperthyroidism; endometriosis; hematopoietic disorders and/or other pathologies and  
 30 disorders. Other GPCR-related diseases and disorders are contemplated.

The protein similarity information, expression pattern, and map location for the  
 olfactory receptor-like GPCR proteins and nucleic acids disclosed herein suggest that these  
 olfactory receptors may have important structural and/or physiological functions  
 characteristic of the olfactory receptor family. Therefore, the GPCR nucleic acids and

proteins are useful in potential diagnostic and therapeutic applications and as a research tool. These include serving as a specific or selective nucleic acid or protein diagnostic and/or prognostic marker, wherein the presence or amount of the nucleic acid or the protein are to be assessed, as well as potential therapeutic applications such as the following: (i) a protein  
 5 therapeutic, (ii) a small molecule drug target, (iii) an antibody target (therapeutic, diagnostic, drug targeting/cytotoxic antibody), (iv) a nucleic acid useful in gene therapy (gene delivery/gene ablation), (v) a composition promoting tissue regeneration in vitro and in vivo, and (vi) a biological defense weapon.

GPCR polypeptides are useful in the generation of antibodies that bind  
 10 immunospecifically to the GPCR polypeptides of the invention, and as vaccines. The antibodies are for use in therapeutic or diagnostic methods. These antibodies may be generated according to methods known in the art, using prediction from hydrophobicity charts, as described in the "Anti-GPCR<sub>X</sub> Antibodies" section below.

GPCR polypeptides can also be used to screen for potential agonist and antagonist  
 15 compounds. For example, a cDNA encoding the GPCR-like protein may be useful in gene therapy, and the GPCR-like protein may be useful when administered to a subject in need thereof. By way of nonlimiting example, the compositions of the present invention will have efficacy for treatment of patients suffering from the diseases and disorders disclosed above and/or other pathologies and disorders. The novel nucleic acid encoding GPCR-like protein,  
 20 and the GPCR-like protein of the invention, or fragments thereof, may further be useful in diagnostic applications, wherein the presence or amount of the nucleic acid or the protein are to be assessed.

### **GPCR<sub>X</sub> Nucleic Acids and Polypeptides**

One aspect of the invention pertains to isolated nucleic acid molecules that encode  
 25 GPCR<sub>X</sub> polypeptides or biologically active portions thereof. Also included in the invention are nucleic acid fragments sufficient for use as hybridization probes to identify GPCR<sub>X</sub>-encoding nucleic acids (*e.g.*, GPCR<sub>X</sub> mRNAs) and fragments for use as PCR primers for the amplification and/or mutation of GPCR<sub>X</sub> nucleic acid molecules. As used herein, the term "nucleic acid molecule" is intended to include DNA molecules (*e.g.*, cDNA or genomic  
 30 DNA), RNA molecules (*e.g.*, mRNA), analogs of the DNA or RNA generated using nucleotide analogs, and derivatives, fragments and homologs thereof. The nucleic acid molecule may be single-stranded or double-stranded, but preferably is comprised double-stranded DNA.

An GPCR<sub>X</sub> nucleic acid can encode a mature GPCR<sub>X</sub> polypeptide. As used herein, a “mature” form of a polypeptide or protein disclosed in the present invention is the product of a naturally occurring polypeptide or precursor form or proprotein. The naturally occurring polypeptide, precursor or proprotein includes, by way of nonlimiting example, the full-length gene product, encoded by the corresponding gene. Alternatively, it may be defined as the polypeptide, precursor or proprotein encoded by an ORF described herein. The product “mature” form arises, again by way of nonlimiting example, as a result of one or more naturally occurring processing steps as they may take place within the cell, or host cell, in which the gene product arises. Examples of such processing steps leading to a “mature” form of a polypeptide or protein include the cleavage of the N-terminal methionine residue encoded by the initiation codon of an ORF, or the proteolytic cleavage of a signal peptide or leader sequence. Thus a mature form arising from a precursor polypeptide or protein that has residues 1 to N, where residue 1 is the N-terminal methionine, would have residues 2 through N remaining after removal of the N-terminal methionine. Alternatively, a mature form arising from a precursor polypeptide or protein having residues 1 to N, in which an N-terminal signal sequence from residue 1 to residue M is cleaved, would have the residues from residue M+1 to residue N remaining. Further as used herein, a “mature” form of a polypeptide or protein may arise from a step of post-translational modification other than a proteolytic cleavage event. Such additional processes include, by way of non-limiting example, glycosylation, myristoylation or phosphorylation. In general, a mature polypeptide or protein may result from the operation of only one of these processes, or a combination of any of them.

The term “probes”, as utilized herein, refers to nucleic acid sequences of variable length, preferably between at least about 10 nucleotides (nt), 100 nt, or as many as approximately, *e.g.*, 6,000 nt, depending upon the specific use. Probes are used in the detection of identical, similar, or complementary nucleic acid sequences. Longer length probes are generally obtained from a natural or recombinant source, are highly specific, and much slower to hybridize than shorter-length oligomer probes. Probes may be single- or double-stranded and designed to have specificity in PCR, membrane-based hybridization technologies, or ELISA-like technologies.

The term “isolated” nucleic acid molecule, as utilized herein, is one, which is separated from other nucleic acid molecules which are present in the natural source of the nucleic acid. Preferably, an “isolated” nucleic acid is free of sequences which naturally flank the nucleic acid (*i.e.*, sequences located at the 5'- and 3'-termini of the nucleic acid) in the

genomic DNA of the organism from which the nucleic acid is derived. For example, in various embodiments, the isolated GPCR<sub>X</sub> nucleic acid molecules can contain less than about 5 kb, 4 kb, 3 kb, 2 kb, 1 kb, 0.5 kb, or 0.1 kb of nucleotide sequences which naturally flank the nucleic acid molecule in genomic DNA of the cell/tissue from which the nucleic acid is derived (*e.g.*, brain, heart, liver, spleen, etc.). Moreover, an "isolated" nucleic acid molecule, such as a cDNA molecule, can be substantially free of other cellular material or culture medium when produced by recombinant techniques, or of chemical precursors or other chemicals when chemically synthesized.

A nucleic acid molecule of the invention, *e.g.*, a nucleic acid molecule having the nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 or a complement of this aforementioned nucleotide sequence, can be isolated using standard molecular biology techniques and the sequence information provided herein. Using all or a portion of the nucleic acid sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 as a hybridization probe, GPCR<sub>X</sub> molecules can be isolated using standard hybridization and cloning techniques (*e.g.*, as described in Sambrook, *et al.*, (eds.), MOLECULAR CLONING: A LABORATORY MANUAL 2<sup>nd</sup> Ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989; and Ausubel, *et al.*, (eds.), CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, New York, NY, 1993.)

A nucleic acid of the invention can be amplified using cDNA, mRNA or alternatively, genomic DNA, as a template and appropriate oligonucleotide primers according to standard PCR amplification techniques. The nucleic acid so amplified can be cloned into an appropriate vector and characterized by DNA sequence analysis. Furthermore, oligonucleotides corresponding to GPCR<sub>X</sub> nucleotide sequences can be prepared by standard synthetic techniques, *e.g.*, using an automated DNA synthesizer.

As used herein, the term "oligonucleotide" refers to a series of linked nucleotide residues, which oligonucleotide has a sufficient number of nucleotide bases to be used in a PCR reaction. A short oligonucleotide sequence may be based on, or designed from, a genomic or cDNA sequence and is used to amplify, confirm, or reveal the presence of an identical, similar or complementary DNA or RNA in a particular cell or tissue. Oligonucleotides comprise portions of a nucleic acid sequence having about 10 nt, 50 nt, or 100 nt in length, preferably about 15 nt to 30 nt in length. In one embodiment of the invention, an oligonucleotide comprising a nucleic acid molecule less than 100 nt in length would further comprise at least 6 contiguous nucleotides of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13,

15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or a complement thereof.

Oligonucleotides may be chemically synthesized and may also be used as probes.

In another embodiment, an isolated nucleic acid molecule of the invention comprises a nucleic acid molecule that is a complement of the nucleotide sequence shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or a portion of this nucleotide sequence (*e.g.*, a fragment that can be used as a probe or primer or a fragment encoding a biologically-active portion of an GPCR<sub>X</sub> polypeptide). A nucleic acid molecule that is complementary to the nucleotide sequence shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 is one that is sufficiently complementary to the nucleotide sequence shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 that it can hydrogen bond with few mismatches to the nucleotide sequence shown SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, thereby forming a stable duplex.

As used herein, the term “complementary” refers to Watson-Crick or Hoogsteen base pairing between nucleotides units of a nucleic acid molecule, and the term “binding” means the physical or chemical interaction between two polypeptides or compounds or associated polypeptides or compounds or combinations thereof. Binding includes ionic, non-ionic, van der Waals, hydrophobic interactions, and the like. A physical interaction can be either direct or indirect. Indirect interactions may be through or due to the effects of another polypeptide or compound. Direct binding refers to interactions that do not take place through, or due to, the effect of another polypeptide or compound, but instead are without other substantial chemical intermediates.

Fragments provided herein are defined as sequences of at least 6 (contiguous) nucleic acids or at least 4 (contiguous) amino acids, a length sufficient to allow for specific hybridization in the case of nucleic acids or for specific recognition of an epitope in the case of amino acids, respectively, and are at most some portion less than a full length sequence. Fragments may be derived from any contiguous portion of a nucleic acid or amino acid sequence of choice. Derivatives are nucleic acid sequences or amino acid sequences formed from the native compounds either directly or by modification or partial substitution. Analogs are nucleic acid sequences or amino acid sequences that have a structure similar to, but not identical to, the native compound but differs from it in respect to certain components or side chains. Analogs may be synthetic or from a different evolutionary origin and may have a similar or opposite metabolic activity compared to wild type. Homologs are nucleic acid



sequences or amino acid sequences of a particular gene that are derived from different species.

Derivatives and analogs may be full length or other than full length, if the derivative or analog contains a modified nucleic acid or amino acid, as described below. Derivatives or analogs of the nucleic acids or proteins of the invention include, but are not limited to, molecules comprising regions that are substantially homologous to the nucleic acids or proteins of the invention, in various embodiments, by at least about 70%, 80%, or 95% identity (with a preferred identity of 80-95%) over a nucleic acid or amino acid sequence of identical size or when compared to an aligned sequence in which the alignment is done by a computer homology program known in the art, or whose encoding nucleic acid is capable of hybridizing to the complement of a sequence encoding the aforementioned proteins under stringent, moderately stringent, or low stringent conditions. See *e.g.* Ausubel, *et al.*, CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, New York, NY, 1993, and below.

A “homologous nucleic acid sequence” or “homologous amino acid sequence,” or variations thereof, refer to sequences characterized by a homology at the nucleotide level or amino acid level as discussed above. Homologous nucleotide sequences encode those sequences coding for isoforms of GPCR<sub>X</sub> polypeptides. Isoforms can be expressed in different tissues of the same organism as a result of, for example, alternative splicing of RNA. Alternatively, isoforms can be encoded by different genes. In the invention, homologous nucleotide sequences include nucleotide sequences encoding for an GPCR<sub>X</sub> polypeptide of species other than humans, including, but not limited to: vertebrates, and thus can include, *e.g.*, frog, mouse, rat, rabbit, dog, cat, cow, horse, and other organisms. Homologous nucleotide sequences also include, but are not limited to, naturally occurring allelic variations and mutations of the nucleotide sequences set forth herein. A homologous nucleotide sequence does not, however, include the exact nucleotide sequence encoding human GPCR<sub>X</sub> protein. Homologous nucleic acid sequences include those nucleic acid sequences that encode conservative amino acid substitutions (see below) in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, as well as a polypeptide possessing GPCR<sub>X</sub> biological activity. Various biological activities of the GPCR<sub>X</sub> proteins are described below.

An GPCR<sub>X</sub> polypeptide is encoded by the open reading frame (“ORF”) of an GPCR<sub>X</sub> nucleic acid. An ORF corresponds to a nucleotide sequence that could potentially be translated into a polypeptide. A stretch of nucleic acids comprising an ORF is uninterrupted

by a stop codon. An ORF that represents the coding sequence for a full protein begins with an ATG "start" codon and terminates with one of the three "stop" codons, namely, TAA, TAG, or TGA. For the purposes of this invention, an ORF may be any part of a coding sequence, with or without a start codon, a stop codon, or both. For an ORF to be considered  
 5 as a good candidate for coding for a *bona fide* cellular protein, a minimum size requirement is often set, *e.g.*, a stretch of DNA that would encode a protein of 50 amino acids or more.

The nucleotide sequences determined from the cloning of the human GPCR<sub>X</sub> genes allows for the generation of probes and primers designed for use in identifying and/or cloning GPCR<sub>X</sub> homologues in other cell types, *e.g.* from other tissues, as well as GPCR<sub>X</sub>  
 10 homologues from other vertebrates. The probe/primer typically comprises substantially purified oligonucleotide. The oligonucleotide typically comprises a region of nucleotide sequence that hybridizes under stringent conditions to an at least about 12, 25, 50, 100, 150, 200, 250, 300, 350 or 400 consecutive sense strand nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39; or an anti-sense strand  
 15 nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39; or of a naturally occurring mutant of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39.

Probes based on the human GPCR<sub>X</sub> nucleotide sequences can be used to detect transcripts or genomic sequences encoding the same or homologous proteins. In various  
 20 embodiments, the probe further comprises a label group attached thereto, *e.g.* the label group can be a radioisotope, a fluorescent compound, an enzyme, or an enzyme co-factor. Such probes can be used as a part of a diagnostic test kit for identifying cells or tissues which mis-express an GPCR<sub>X</sub> protein, such as by measuring a level of an GPCR<sub>X</sub>-encoding nucleic acid in a sample of cells from a subject *e.g.*, detecting GPCR<sub>X</sub> mRNA levels or determining  
 25 whether a genomic GPCR<sub>X</sub> gene has been mutated or deleted.

"A polypeptide having a biologically-active portion of an GPCR<sub>X</sub> polypeptide" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. A nucleic acid fragment encoding a "biologically-  
 30 active portion of GPCR<sub>X</sub>" can be prepared by isolating a portion SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 that encodes a polypeptide having an GPCR<sub>X</sub> biological activity (the biological activities of the GPCR<sub>X</sub> proteins are described below), expressing the encoded portion of GPCR<sub>X</sub> protein (*e.g.*, by recombinant expression *in vitro*) and assessing the activity of the encoded portion of GPCR<sub>X</sub>.

## GPCRX Nucleic Acid and Polypeptide Variants

The invention further encompasses nucleic acid molecules that differ from the nucleotide sequences shown SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 due to degeneracy of the genetic code and thus encode the same GPCR proteins as that encoded by the nucleotide sequences shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39. In another embodiment, an isolated nucleic acid molecule of the invention has a nucleotide sequence encoding a protein having an amino acid sequence shown in SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40. In addition to the human GPCR nucleotide sequences shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, it will be appreciated by those skilled in the art that DNA sequence polymorphisms that lead to changes in the amino acid sequences of the GPCR polypeptides may exist within a population (*e.g.*, the human population). Such genetic polymorphism in the GPCR genes may exist among individuals within a population due to natural allelic variation. As used herein, the terms "gene" and "recombinant gene" refer to nucleic acid molecules comprising an open reading frame (ORF) encoding an GPCR protein, preferably a vertebrate GPCR protein. Such natural allelic variations can typically result in 1-5% variance in the nucleotide sequence of the GPCR genes. Any and all such nucleotide variations and resulting amino acid polymorphisms in the GPCR polypeptides, which are the result of natural allelic variation and that do not alter the functional activity of the GPCR polypeptides, are intended to be within the scope of the invention.

Moreover, nucleic acid molecules encoding GPCR proteins from other species, and thus that have a nucleotide sequence that differs from the human sequence SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 are intended to be within the scope of the invention. Nucleic acid molecules corresponding to natural allelic variants and homologues of the GPCR cDNAs of the invention can be isolated based on their homology to the human GPCR nucleic acids disclosed herein using the human cDNAs, or a portion thereof, as a hybridization probe according to standard hybridization techniques under stringent hybridization conditions.

Accordingly, in another embodiment, an isolated nucleic acid molecule of the invention is at least 6 nucleotides in length and hybridizes under stringent conditions to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39.

In another embodiment, the nucleic acid is at least 10, 25, 50, 100, 250, 500, 750, 1000, 1500, or 2000 or more nucleotides in length. In yet another embodiment, an isolated nucleic acid molecule of the invention hybridizes to the coding region. As used herein, the term "hybridizes under stringent conditions" is intended to describe conditions for hybridization and washing under which nucleotide sequences at least 60% homologous to each other typically remain hybridized to each other.

Homologs (*i.e.*, nucleic acids encoding GPCR<sub>X</sub> proteins derived from species other than human) or other related sequences (*e.g.*, paralogs) can be obtained by low, moderate or high stringency hybridization with all or a portion of the particular human sequence as a probe using methods well known in the art for nucleic acid hybridization and cloning.

As used herein, the phrase "stringent hybridization conditions" refers to conditions under which a probe, primer or oligonucleotide will hybridize to its target sequence, but to no other sequences. Stringent conditions are sequence-dependent and will be different in different circumstances. Longer sequences hybridize specifically at higher temperatures than shorter sequences. Generally, stringent conditions are selected to be about 5 °C lower than the thermal melting point (*T<sub>m</sub>*) for the specific sequence at a defined ionic strength and pH. The *T<sub>m</sub>* is the temperature (under defined ionic strength, pH and nucleic acid concentration) at which 50% of the probes complementary to the target sequence hybridize to the target sequence at equilibrium. Since the target sequences are generally present at excess, at *T<sub>m</sub>*, 50% of the probes are occupied at equilibrium. Typically, stringent conditions will be those in which the salt concentration is less than about 1.0 M sodium ion, typically about 0.01 to 1.0 M sodium ion (or other salts) at pH 7.0 to 8.3 and the temperature is at least about 30 °C for short probes, primers or oligonucleotides (*e.g.*, 10 nt to 50 nt) and at least about 60 °C for longer probes, primers and oligonucleotides. Stringent conditions may also be achieved with the addition of destabilizing agents, such as formamide.

Stringent conditions are known to those skilled in the art and can be found in Ausubel, *et al.*, (eds.), CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. Preferably, the conditions are such that sequences at least about 65%, 70%, 75%, 85%, 90%, 95%, 98%, or 99% homologous to each other typically remain hybridized to each other. A non-limiting example of stringent hybridization conditions are hybridization in a high salt buffer comprising 6X SSC, 50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 0.02% PVP, 0.02% Ficoll, 0.02% BSA, and 500 mg/ml denatured salmon sperm DNA at 65 °C, followed by one or more washes in 0.2X SSC, 0.01% BSA at 50 °C. An isolated nucleic acid molecule of the invention that hybridizes under stringent conditions to

the sequences of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 corresponds to a naturally-occurring nucleic acid molecule. As used herein, a "naturally-occurring" nucleic acid molecule refers to an RNA or DNA molecule having a nucleotide sequence that occurs in nature (*e.g.*, encodes a natural protein).

5 In a second embodiment, a nucleic acid sequence that is hybridizable to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 or fragments, analogs or derivatives thereof, under conditions of moderate stringency is provided. A non-limiting example of moderate stringency hybridization conditions are hybridization in 6X SSC, 5X Denhardt's solution,  
10 0.5% SDS and 100 mg/ml denatured salmon sperm DNA at 55 °C, followed by one or more washes in 1X SSC, 0.1% SDS at 37 °C. Other conditions of moderate stringency that may be used are well-known within the art. See, *e.g.*, Ausubel, *et al.* (eds.), 1993, CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, NY, and Kriegler, 1990; GENE TRANSFER AND EXPRESSION, A LABORATORY MANUAL, Stockton Press, NY.

15 In a third embodiment, a nucleic acid that is hybridizable to the nucleic acid molecule comprising the nucleotide sequences of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 or fragments, analogs or derivatives thereof, under conditions of low stringency, is provided. A non-limiting example of low stringency hybridization conditions are hybridization in 35% formamide, 5X SSC, 50 mM Tris-HCl (pH  
20 7.5), 5 mM EDTA, 0.02% PVP, 0.02% Ficoll, 0.2% BSA, 100 mg/ml denatured salmon sperm DNA, 10% (wt/vol) dextran sulfate at 40 °C, followed by one or more washes in 2X SSC, 25 mM Tris-HCl (pH 7.4), 5 mM EDTA, and 0.1% SDS at 50 °C. Other conditions of low stringency that may be used are well known in the art (*e.g.*, as employed for cross-species hybridizations). See, *e.g.*, Ausubel, *et al.* (eds.), 1993, CURRENT PROTOCOLS IN  
25 MOLECULAR BIOLOGY, John Wiley & Sons, NY, and Kriegler, 1990, GENE TRANSFER AND EXPRESSION, A LABORATORY MANUAL, Stockton Press, NY; Shilo and Weinberg, 1981. *Proc Natl Acad Sci USA* 78: 6789-6792.

### Conservative Mutations

30 In addition to naturally-occurring allelic variants of GPCR<sub>X</sub> sequences that may exist in the population, the skilled artisan will further appreciate that changes can be introduced by mutation into the nucleotide sequences of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 thereby leading to changes in the amino acid sequences of the encoded GPCR<sub>X</sub> proteins, without altering the functional ability of said GPCR<sub>X</sub>

proteins. For example, nucleotide substitutions leading to amino acid substitutions at "non-essential" amino acid residues can be made in the sequence of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40. A "non-essential" amino acid residue is a residue that can be altered from the wild-type sequences of the GPCR<sub>X</sub> proteins without altering their biological activity, whereas an "essential" amino acid residue is required for such biological activity. For example, amino acid residues that are conserved among the GPCR<sub>X</sub> proteins of the invention are predicted to be particularly non-amenable to alteration. Amino acids for which conservative substitutions can be made are well-known within the art.

Another aspect of the invention pertains to nucleic acid molecules encoding GPCR<sub>X</sub> proteins that contain changes in amino acid residues that are not essential for activity. Such GPCR<sub>X</sub> proteins differ in amino acid sequence from SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40 yet retain biological activity. In one embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a protein, wherein the protein comprises an amino acid sequence at least about 45% homologous to the amino acid sequences of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40. Preferably, the protein encoded by the nucleic acid molecule is at least about 60% homologous to SEQ ID NOS: SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40; more preferably at least about 70% homologous to SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40; still more preferably at least about 80% homologous to SEQ ID NOS: SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40; even more preferably at least about 90% homologous to SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40; and most preferably at least about 95% homologous to SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40.

An isolated nucleic acid molecule encoding an GPCR<sub>X</sub> protein homologous to the protein of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40 can be created by introducing one or more nucleotide substitutions, additions or deletions into the nucleotide sequence of SEQ ID NOS: 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 such that one or more amino acid substitutions, additions or deletions are introduced into the encoded protein.

Mutations can be introduced into SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40 by standard techniques, such as site-directed mutagenesis and PCR-mediated mutagenesis. Preferably, conservative amino acid substitutions are made

at one or more predicted, non-essential amino acid residues. A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined within the art. These families include amino acids with basic side chains (*e.g.*, lysine, arginine, histidine), acidic side chains (*e.g.*, aspartic acid, glutamic acid), uncharged polar side chains (*e.g.*, glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (*e.g.*, alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), beta-branched side chains (*e.g.*, threonine, valine, isoleucine) and aromatic side chains (*e.g.*, tyrosine, phenylalanine, tryptophan, histidine). Thus, a predicted non-essential amino acid residue in the GPCR<sub>X</sub> protein is replaced with another amino acid residue from the same side chain family. Alternatively, in another embodiment, mutations can be introduced randomly along all or part of an GPCR<sub>X</sub> coding sequence, such as by saturation mutagenesis, and the resultant mutants can be screened for GPCR<sub>X</sub> biological activity to identify mutants that retain activity. Following mutagenesis of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, the encoded protein can be expressed by any recombinant technology known in the art and the activity of the protein can be determined.

The relatedness of amino acid families may also be determined based on side chain interactions. Substituted amino acids may be fully conserved "strong" residues or fully conserved "weak" residues. The "strong" group of conserved amino acid residues may be any one of the following groups: STA, NEQK, NHQK, NDEQ, QHRK, MILV, MILF, HY, FYW, wherein the single letter amino acid codes are grouped by those amino acids that may be substituted for each other. Likewise, the "weak" group of conserved residues may be any one of the following: CSA, ATV, SAG, STNK, STPA, SGND, SNDEQK, NDEQHK, NEQHRK, VLIM, HFY, wherein the letters within each group represent the single letter amino acid code.

In one embodiment, a mutant GPCR<sub>X</sub> protein can be assayed for (i) the ability to form protein:protein interactions with other GPCR<sub>X</sub> proteins, other cell-surface proteins, or biologically-active portions thereof, (ii) complex formation between a mutant GPCR<sub>X</sub> protein and an GPCR<sub>X</sub> ligand; or (iii) the ability of a mutant GPCR<sub>X</sub> protein to bind to an intracellular target protein or biologically-active portion thereof.

In yet another embodiment, a mutant GPCR<sub>X</sub> protein can be assayed for the ability to regulate a specific biological function (*e.g.*, regulation of insulin release).

## Antisense Nucleic Acids

Another aspect of the invention pertains to isolated antisense nucleic acid molecules that are hybridizable to or complementary to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or fragments, analogs or derivatives thereof. An "antisense" nucleic acid comprises a nucleotide sequence that is complementary to a "sense" nucleic acid encoding a protein (*e.g.*, complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence). In specific aspects, antisense nucleic acid molecules are provided that comprise a sequence complementary to at least about 10, 25, 50, 100, 250 or 500 nucleotides or an entire GPCR<sub>X</sub> coding strand, or to only a portion thereof. Nucleic acid molecules encoding fragments, homologs, derivatives and analogs of an GPCR<sub>X</sub> protein of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or antisense nucleic acids complementary to an GPCR<sub>X</sub> nucleic acid sequence of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, are additionally provided.

In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence encoding an GPCR<sub>X</sub> protein. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues. In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence encoding the GPCR<sub>X</sub> protein. The term "noncoding region" refers to 5' and 3' sequences which flank the coding region that are not translated into amino acids (*i.e.*, also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding the GPCR<sub>X</sub> protein disclosed herein, antisense nucleic acids of the invention can be designed according to the rules of Watson and Crick or Hoogsteen base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of GPCR<sub>X</sub> mRNA, but more preferably is an oligonucleotide that is antisense to only a portion of the coding or noncoding region of GPCR<sub>X</sub> mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of GPCR<sub>X</sub> mRNA. An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis or enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (*e.g.*, an antisense oligonucleotide) can be chemically synthesized using naturally-occurring nucleotides or



variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids (*e.g.*, phosphorothioate derivatives and acridine substituted nucleotides can be used).

5           Examples of modified nucleotides that can be used to generate the antisense nucleic acid include: 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 10   2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 15   5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (*i.e.*, RNA transcribed from the inserted nucleic acid will be of an antisense 20   orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated *in situ* such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding an GPCR protein to thereby inhibit expression of the protein (*e.g.*, by inhibiting transcription and/or translation). The hybridization can be by conventional 25   nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule that binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site.

Alternatively, antisense nucleic acid molecules can be modified to target selected cells 30   and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface (*e.g.*, by linking the antisense nucleic acid molecules to peptides or antibodies that bind to cell surface receptors or antigens). The antisense nucleic acid molecules can also be delivered to cells using the vectors described herein. To achieve

sufficient nucleic acid molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an  $\alpha$ -anomeric nucleic acid molecule. An  $\alpha$ -anomeric nucleic acid molecule forms specific  
 5 double-stranded hybrids with complementary RNA in which, contrary to the usual  $\beta$ -units, the strands run parallel to each other. See, *e.g.*, Gaultier, *et al.*, 1987. *Nucl. Acids Res.* **15**: 6625-6641. The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (see, *e.g.*, Inoue, *et al.* 1987. *Nucl. Acids Res.* **15**: 6131-6148) or a chimeric RNA-DNA analogue (see, *e.g.*, Inoue, *et al.*, 1987. *FEBS Lett.* **215**: 327-330.

10

### **Ribozymes and PNA Moieties**

Nucleic acid modifications include, by way of non-limiting example, modified bases, and nucleic acids whose sugar phosphate backbones are modified or derivatized. These modifications are carried out at least in part to enhance the chemical stability of the modified  
 15 nucleic acid, such that they may be used, for example, as antisense binding nucleic acids in therapeutic applications in a subject.

In one embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity that are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a  
 20 complementary region. Thus, ribozymes (*e.g.*, hammerhead ribozymes as described in Haselhoff and Gerlach 1988. *Nature* 334: 585-591) can be used to catalytically cleave GPCR<sub>X</sub> mRNA transcripts to thereby inhibit translation of GPCR<sub>X</sub> mRNA. A ribozyme having specificity for an GPCR<sub>X</sub>-encoding nucleic acid can be designed based upon the nucleotide sequence of an GPCR<sub>X</sub> cDNA disclosed herein (*i.e.*, SEQ ID NOS:1, 3, 5, 7, 9,  
 25 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39). For example, a derivative of a *Tetrahymena* L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in an GPCR<sub>X</sub>-encoding mRNA. See, *e.g.*, U.S. Patent 4,987,071 to Cech, *et al.* and U.S. Patent 5,116,742 to Cech, *et al.* GPCR<sub>X</sub> mRNA can also be used to select a catalytic RNA having a specific ribonuclease  
 30 activity from a pool of RNA molecules. See, *e.g.*, Bartel *et al.*, (1993) *Science* 261:1411-1418.

Alternatively, GPCR<sub>X</sub> gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region of the GPCR<sub>X</sub> nucleic acid (*e.g.*, the GPCR<sub>X</sub> promoter and/or enhancers) to form triple helical structures that prevent transcription

of the GPCR<sub>X</sub> gene in target cells. *See, e.g., Helene, 1991. Anticancer Drug Des. 6: 569-84; Helene, et al. 1992. Ann. N.Y. Acad. Sci. 660: 27-36; Maher, 1992. Bioassays 14: 807-15.*

In various embodiments, the GPCR<sub>X</sub> nucleic acids can be modified at the base moiety, sugar moiety or phosphate backbone to improve, *e.g.,* the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids. *See, e.g., Hyrup, et al., 1996. Bioorg Med Chem 4: 5-23.* As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics (*e.g.,* DNA mimics) in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained.

The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength. The synthesis of PNA oligomers can be performed using standard solid phase peptide synthesis protocols as described in Hyrup, *et al., 1996, supra; Perry-O'Keefe, et al., 1996. Proc. Natl. Acad. Sci. USA 93: 14670-14675.*

PNAs of GPCR<sub>X</sub> can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, *e.g.,* inducing transcription or translation arrest or inhibiting replication. PNAs of GPCR<sub>X</sub> can also be used, for example, in the analysis of single base pair mutations in a gene (*e.g.,* PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, *e.g.,* S<sub>1</sub> nucleases (*see, Hyrup, et al., 1996, supra; Perry-O'Keefe, et al., 1996, supra*); or as probes or primers for DNA sequence and hybridization (*see, Hyrup, et al., 1996, supra; Perry-O'Keefe, et al., 1996, supra*).

In another embodiment, PNAs of GPCR<sub>X</sub> can be modified, *e.g.,* to enhance their stability or cellular uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras of GPCR<sub>X</sub> can be generated that may combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes (*e.g.,* RNase H and DNA polymerases) to interact with the DNA portion while the PNA portion would provide high binding affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths selected in terms of base stacking, number of bonds between the nucleobases, and orientation (*see, Hyrup, et al., 1996, supra*). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup, *et al., 1996, supra* and Finn, *et al., 1996. Nucl Acids Res 24: 3357-3363.* For example, a DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry, and modified nucleoside analogs, *e.g.,* 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine

phosphoramidite, can be used between the PNA and the 5' end of DNA. *See, e.g., Mag, et al., 1989 Nucl Acid Res 17: 5973-5988.* PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment. *See, e.g., Finn, et al., 1996, supra.* Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment. *See, e.g., Petersen, et al., 1975 Bioorg. Med. Chem. Lett. 5: 1119-11124.*

In other embodiments, the oligonucleotide may include other appended groups such as peptides (*e.g., for targeting host cell receptors in vivo*), or agents facilitating transport across the cell membrane (*see, e.g., Letsinger, et al., 1989 Proc. Natl. Acad. Sci. U.S.A. 86: 6553-6556; Lemaitre, et al., 1987 Proc. Natl. Acad. Sci. 84: 648-652; PCT Publication No. WO88/09810*) or the blood-brain barrier (*see, e.g., PCT Publication No. WO 89/10134*). In addition, oligonucleotides can be modified with hybridization triggered cleavage agents (*see, e.g., Krol, et al., 1988 BioTechniques 6:958-976*) or intercalating agents (*see, e.g., Zon, 1988 Pharm. Res. 5: 539-549*). To this end, the oligonucleotide may be conjugated to another molecule, *e.g., a peptide, a hybridization triggered cross-linking agent, a transport agent, a hybridization-triggered cleavage agent, and the like.*

### GPCRX Polypeptides

A polypeptide according to the invention includes a polypeptide including the amino acid sequence of GPCR<sub>X</sub> polypeptides whose sequences are provided in SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40. The invention also includes a mutant or variant protein any of whose residues may be changed from the corresponding residues shown in SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40 while still encoding a protein that maintains its GPCR<sub>X</sub> activities and physiological functions, or a functional fragment thereof.

In general, an GPCR<sub>X</sub> variant that preserves GPCR<sub>X</sub>-like function includes any variant in which residues at a particular position in the sequence have been substituted by other amino acids, and further include the possibility of inserting an additional residue or residues between two residues of the parent protein as well as the possibility of deleting one or more residues from the parent sequence. Any amino acid substitution, insertion, or deletion is encompassed by the invention. In favorable circumstances, the substitution is a conservative substitution as defined above.

One aspect of the invention pertains to isolated GPCR<sub>X</sub> proteins, and biologically-active portions thereof, or derivatives, fragments, analogs or homologs thereof. Also

provided are polypeptide fragments suitable for use as immunogens to raise anti-GPCR<sub>X</sub> antibodies. In one embodiment, native GPCR<sub>X</sub> proteins can be isolated from cells or tissue sources by an appropriate purification scheme using standard protein purification techniques. In another embodiment, GPCR<sub>X</sub> proteins are produced by recombinant DNA techniques.

5 Alternative to recombinant expression, an GPCR<sub>X</sub> protein or polypeptide can be synthesized chemically using standard peptide synthesis techniques.

An "isolated" or "purified" polypeptide or protein or biologically-active portion thereof is substantially free of cellular material or other contaminating proteins from the cell or tissue source from which the GPCR<sub>X</sub> protein is derived, or substantially free from  
10 chemical precursors or other chemicals when chemically synthesized. The language "substantially free of cellular material" includes preparations of GPCR<sub>X</sub> proteins in which the protein is separated from cellular components of the cells from which it is isolated or recombinantly-produced. In one embodiment, the language "substantially free of cellular material" includes preparations of GPCR<sub>X</sub> proteins having less than about 30% (by dry  
15 weight) of non-GPCR<sub>X</sub> proteins (also referred to herein as a "contaminating protein"), more preferably less than about 20% of non-GPCR<sub>X</sub> proteins, still more preferably less than about 10% of non-GPCR<sub>X</sub> proteins, and most preferably less than about 5% of non-GPCR<sub>X</sub> proteins. When the GPCR<sub>X</sub> protein or biologically-active portion thereof is recombinantly-produced, it is also preferably substantially free of culture medium, *i.e.*, culture medium  
20 represents less than about 20%, more preferably less than about 10%, and most preferably less than about 5% of the volume of the GPCR<sub>X</sub> protein preparation.

The language "substantially free of chemical precursors or other chemicals" includes preparations of GPCR<sub>X</sub> proteins in which the protein is separated from chemical precursors or other chemicals that are involved in the synthesis of the protein. In one embodiment, the  
25 language "substantially free of chemical precursors or other chemicals" includes preparations of GPCR<sub>X</sub> proteins having less than about 30% (by dry weight) of chemical precursors or non-GPCR<sub>X</sub> chemicals, more preferably less than about 20% chemical precursors or non-GPCR<sub>X</sub> chemicals, still more preferably less than about 10% chemical precursors or non-GPCR<sub>X</sub> chemicals, and most preferably less than about 5% chemical precursors or  
30 non-GPCR<sub>X</sub> chemicals.

Biologically-active portions of GPCR<sub>X</sub> proteins include peptides comprising amino acid sequences sufficiently homologous to or derived from the amino acid sequences of the GPCR<sub>X</sub> proteins (*e.g.*, the amino acid sequence shown in SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40) that include fewer amino acids than the

full-length GPCR<sub>X</sub> proteins, and exhibit at least one activity of an GPCR<sub>X</sub> protein.

Typically, biologically-active portions comprise a domain or motif with at least one activity of the GPCR<sub>X</sub> protein. A biologically-active portion of an GPCR<sub>X</sub> protein can be a polypeptide which is, for example, 10, 25, 50, 100 or more amino acid residues in length.

5           Moreover, other biologically-active portions, in which other regions of the protein are deleted, can be prepared by recombinant techniques and evaluated for one or more of the functional activities of a native GPCR<sub>X</sub> protein.

          In an embodiment, the GPCR<sub>X</sub> protein has an amino acid sequence shown in SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40. In other  
10       embodiments, the GPCR<sub>X</sub> protein is substantially homologous to SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 34, 36, 38 and 40, and retains the functional activity of the protein of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40, yet differs in amino acid sequence due to natural allelic variation or mutagenesis, as described in detail, below. Accordingly, in another embodiment, the GPCR<sub>X</sub> protein is a  
15       protein that comprises an amino acid sequence at least about 45% homologous to the amino acid sequence SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40, and retains the functional activity of the GPCR<sub>X</sub> proteins of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40.

## 20       **Determining Homology Between Two or More Sequences**

          To determine the percent homology of two amino acid sequences or of two nucleic acids, the sequences are aligned for optimal comparison purposes (*e.g.*, gaps can be introduced in the sequence of a first amino acid or nucleic acid sequence for optimal alignment with a second amino or nucleic acid sequence). The amino acid residues or  
25       nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, then the molecules are homologous at that position (*i.e.*, as used herein amino acid or nucleic acid "homology" is equivalent to amino acid or nucleic acid "identity").

30       The nucleic acid sequence homology may be determined as the degree of identity between two sequences. The homology may be determined using computer programs known in the art, such as GAP software provided in the GCG program package. *See*, Needleman and Wunsch, 1970 *J Mol Biol* 48: 443-453. Using GCG GAP software with the following settings for nucleic acid sequence comparison: GAP creation penalty of 5.0 and GAP

extension penalty of 0.3, the coding region of the analogous nucleic acid sequences referred to above exhibits a degree of identity preferably of at least 70%, 75%, 80%, 85%, 90%, 95%, 98%, or 99%, with the CDS (encoding) part of the DNA sequence shown in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39.

5           The term "sequence identity" refers to the degree to which two polynucleotide or polypeptide sequences are identical on a residue-by-residue basis over a particular region of comparison. The term "percentage of sequence identity" is calculated by comparing two optimally aligned sequences over that region of comparison, determining the number of positions at which the identical nucleic acid base (*e.g.*, A, T, C, G, U, or I, in the case of  
10           nucleic acids) occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the region of comparison (*i.e.*, the window size), and multiplying the result by 100 to yield the percentage of sequence identity. The term "substantial identity" as used herein denotes a characteristic of a polynucleotide sequence, wherein the polynucleotide comprises a sequence that has at least  
15           80 percent sequence identity, preferably at least 85 percent identity and often 90 to 95 percent sequence identity, more usually at least 99 percent sequence identity as compared to a reference sequence over a comparison region.

### **Chimeric and Fusion Proteins**

20           The invention also provides GPCR<sub>X</sub> chimeric or fusion proteins. As used herein, an GPCR<sub>X</sub> "chimeric protein" or "fusion protein" comprises an GPCR<sub>X</sub> polypeptide operatively-linked to a non-GPCR<sub>X</sub> polypeptide. An "GPCR<sub>X</sub> polypeptide" refers to a polypeptide having an amino acid sequence corresponding to an GPCR<sub>X</sub> protein (SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36, 38 and 40), whereas a  
25           "non-GPCR<sub>X</sub> polypeptide" refers to a polypeptide having an amino acid sequence corresponding to a protein that is not substantially homologous to the GPCR<sub>X</sub> protein, *e.g.*, a protein that is different from the GPCR<sub>X</sub> protein and that is derived from the same or a different organism. Within an GPCR<sub>X</sub> fusion protein the GPCR<sub>X</sub> polypeptide can correspond to all or a portion of an GPCR<sub>X</sub> protein. In one embodiment, an GPCR<sub>X</sub> fusion  
30           protein comprises at least one biologically-active portion of an GPCR<sub>X</sub> protein. In another embodiment, an GPCR<sub>X</sub> fusion protein comprises at least two biologically-active portions of an GPCR<sub>X</sub> protein. In yet another embodiment, an GPCR<sub>X</sub> fusion protein comprises at least three biologically-active portions of an GPCR<sub>X</sub> protein. Within the fusion protein, the term "operatively-linked" is intended to indicate that the GPCR<sub>X</sub> polypeptide and the non-GPCR<sub>X</sub>

polypeptide are fused in-frame with one another. The non-GPCRX polypeptide can be fused to the N-terminus or C-terminus of the GPCRX polypeptide.

In one embodiment, the fusion protein is a GST-GPCRX fusion protein in which the GPCRX sequences are fused to the C-terminus of the GST (glutathione S-transferase) sequences. Such fusion proteins can facilitate the purification of recombinant GPCRX polypeptides.

In another embodiment, the fusion protein is an GPCRX protein containing a heterologous signal sequence at its N-terminus. In certain host cells (*e.g.*, mammalian host cells), expression and/or secretion of GPCRX can be increased through use of a heterologous signal sequence.

In yet another embodiment, the fusion protein is an GPCRX-immunoglobulin fusion protein in which the GPCRX sequences are fused to sequences derived from a member of the immunoglobulin protein family. The GPCRX-immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between an GPCRX ligand and an GPCRX protein on the surface of a cell, to thereby suppress GPCRX-mediated signal transduction *in vivo*. The GPCRX-immunoglobulin fusion proteins can be used to affect the bioavailability of an GPCRX cognate ligand. Inhibition of the GPCRX ligand/GPCRX interaction may be useful therapeutically for both the treatment of proliferative and differentiative disorders, as well as modulating (*e.g.* promoting or inhibiting) cell survival. Moreover, the GPCRX-immunoglobulin fusion proteins of the invention can be used as immunogens to produce anti-GPCRX antibodies in a subject, to purify GPCRX ligands, and in screening assays to identify molecules that inhibit the interaction of GPCRX with an GPCRX ligand.

An GPCRX chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, *e.g.*, by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (*see, e.g.*, Ausubel, *et al.* (eds.) CURRENT PROTOCOLS IN



MOLECULAR BIOLOGY, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (*e.g.*, a GST polypeptide). An GPCR<sub>X</sub>-encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the GPCR<sub>X</sub> protein.

5

### GPCR<sub>X</sub> Agonists and Antagonists

The invention also pertains to variants of the GPCR<sub>X</sub> proteins that function as either GPCR<sub>X</sub> agonists (*i.e.*, mimetics) or as GPCR<sub>X</sub> antagonists. Variants of the GPCR<sub>X</sub> protein can be generated by mutagenesis (*e.g.*, discrete point mutation or truncation of the GPCR<sub>X</sub> protein). An agonist of the GPCR<sub>X</sub> protein can retain substantially the same, or a subset of, the biological activities of the naturally occurring form of the GPCR<sub>X</sub> protein. An antagonist of the GPCR<sub>X</sub> protein can inhibit one or more of the activities of the naturally occurring form of the GPCR<sub>X</sub> protein by, for example, competitively binding to a downstream or upstream member of a cellular signaling cascade which includes the GPCR<sub>X</sub> protein. Thus, specific biological effects can be elicited by treatment with a variant of limited function. In one embodiment, treatment of a subject with a variant having a subset of the biological activities of the naturally occurring form of the protein has fewer side effects in a subject relative to treatment with the naturally occurring form of the GPCR<sub>X</sub> proteins.

Variants of the GPCR<sub>X</sub> proteins that function as either GPCR<sub>X</sub> agonists (*i.e.*, mimetics) or as GPCR<sub>X</sub> antagonists can be identified by screening combinatorial libraries of mutants (*e.g.*, truncation mutants) of the GPCR<sub>X</sub> proteins for GPCR<sub>X</sub> protein agonist or antagonist activity. In one embodiment, a variegated library of GPCR<sub>X</sub> variants is generated by combinatorial mutagenesis at the nucleic acid level and is encoded by a variegated gene library. A variegated library of GPCR<sub>X</sub> variants can be produced by, for example, enzymatically ligating a mixture of synthetic oligonucleotides into gene sequences such that a degenerate set of potential GPCR<sub>X</sub> sequences is expressible as individual polypeptides, or alternatively, as a set of larger fusion proteins (*e.g.*, for phage display) containing the set of GPCR<sub>X</sub> sequences therein. There are a variety of methods which can be used to produce libraries of potential GPCR<sub>X</sub> variants from a degenerate oligonucleotide sequence. Chemical synthesis of a degenerate gene sequence can be performed in an automatic DNA synthesizer, and the synthetic gene then ligated into an appropriate expression vector. Use of a degenerate set of genes allows for the provision, in one mixture, of all of the sequences encoding the desired set of potential GPCR<sub>X</sub> sequences. Methods for synthesizing degenerate oligonucleotides are well-known within the art. *See, e.g.*, Narang, 1983 *Tetrahedron* 39:3;

Itakura, *et al.*, 1984 *Annu. Rev. Biochem.* 53:323; Itakura, *et al.*, 1984 *Science* 198:1056; Ike, *et al.*, 1983 *Nucl. Acids Res.* 11:477.

### Polypeptide Libraries

5           In addition, libraries of fragments of the GPCR<sub>X</sub> protein coding sequences can be used to generate a variegated population of GPCR<sub>X</sub> fragments for screening and subsequent selection of variants of an GPCR<sub>X</sub> protein. In one embodiment, a library of coding sequence fragments can be generated by treating a double stranded PCR fragment of an GPCR<sub>X</sub> coding sequence with a nuclease under conditions wherein nicking occurs only about once  
10       per molecule, denaturing the double stranded DNA, renaturing the DNA to form double-stranded DNA that can include sense/antisense pairs from different nicked products, removing single stranded portions from reformed duplexes by treatment with S<sub>1</sub> nuclease, and ligating the resulting fragment library into an expression vector. By this method, expression libraries can be derived which encodes N-terminal and internal fragments of  
15       various sizes of the GPCR<sub>X</sub> proteins.

          Various techniques are known in the art for screening gene products of combinatorial libraries made by point mutations or truncation, and for screening cDNA libraries for gene products having a selected property. Such techniques are adaptable for rapid screening of the gene libraries generated by the combinatorial mutagenesis of GPCR<sub>X</sub> proteins. The most  
20       widely used techniques, which are amenable to high throughput analysis, for screening large gene libraries typically include cloning the gene library into replicable expression vectors, transforming appropriate cells with the resulting library of vectors, and expressing the combinatorial genes under conditions in which detection of a desired activity facilitates isolation of the vector encoding the gene whose product was detected. Recursive ensemble  
25       mutagenesis (REM), a new technique that enhances the frequency of functional mutants in the libraries, can be used in combination with the screening assays to identify GPCR<sub>X</sub> variants. See, *e.g.*, Arkin and Youvan, 1992 *Proc. Natl. Acad. Sci. USA* 89:7811-7815; Delgrave, *et al.*, 1993 *Protein Engineering* 6:327-331.

### Anti-GPCR<sub>X</sub> Antibodies

30           Also included in the invention are antibodies to GPCR<sub>X</sub> proteins, or fragments of GPCR<sub>X</sub> proteins. The term "antibody" as used herein refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, *i.e.*, molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. Such

antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, F<sub>ab</sub>, F<sub>ab</sub>' and F<sub>(ab')<sub>2</sub></sub> fragments, and an F<sub>ab</sub> expression library. In general, an antibody molecule obtained from humans relates to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes  
5 have subclasses as well, such as IgG<sub>1</sub>, IgG<sub>2</sub>, and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain. Reference herein to antibodies includes a reference to all such classes, subclasses and types of human antibody species.

An isolated GPCR<sub>X</sub>-related protein of the invention may be intended to serve as an antigen, or a portion or fragment thereof, and additionally can be used as an immunogen to  
10 generate antibodies that immunospecifically bind the antigen, using standard techniques for polyclonal and monoclonal antibody preparation. The full-length protein can be used or, alternatively, the invention provides antigenic peptide fragments of the antigen for use as immunogens. An antigenic peptide fragment comprises at least 6 amino acid residues of the amino acid sequence of the full length protein and encompasses an epitope thereof such that  
15 an antibody raised against the peptide forms a specific immune complex with the full length protein or with any fragment that contains the epitope. Preferably, the antigenic peptide comprises at least 10 amino acid residues, or at least 15 amino acid residues, or at least 20 amino acid residues, or at least 30 amino acid residues. Preferred epitopes encompassed by the antigenic peptide are regions of the protein that are located on its surface; commonly  
20 these are hydrophilic regions.

In certain embodiments of the invention, at least one epitope encompassed by the antigenic peptide is a region of GPCR<sub>X</sub>-related protein that is located on the surface of the protein, *e.g.*, a hydrophilic region. A hydrophobicity analysis of the human GPCR<sub>X</sub>-related protein sequence will indicate which regions of a GPCR<sub>X</sub>-related protein are particularly  
25 hydrophilic and, therefore, are likely to encode surface residues useful for targeting antibody production. As a means for targeting antibody production, hydropathy plots showing regions of hydrophilicity and hydrophobicity may be generated by any method well known in the art, including, for example, the Kyte Doolittle or the Hopp Woods methods, either with or without Fourier transformation. See, *e.g.*, Hopp and Woods, 1981, *Proc. Nat. Acad. Sci. USA*  
30 78: 3824-3828; Kyte and Doolittle 1982, *J. Mol. Biol.* 157: 105-142, each of which is incorporated herein by reference in its entirety. Antibodies that are specific for one or more domains within an antigenic protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

A protein of the invention, or a derivative, fragment, analog, homolog or ortholog thereof, may be utilized as an immunogen in the generation of antibodies that immunospecifically bind these protein components.

Various procedures known within the art may be used for the production of polyclonal or monoclonal antibodies directed against a protein of the invention, or against derivatives, fragments, analogs homologs or orthologs thereof (see, for example, Antibodies: A Laboratory Manual, Harlow and Lane, 1988, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, incorporated herein by reference). Some of these antibodies are discussed below.

### **Polyclonal Antibodies**

For the production of polyclonal antibodies, various suitable host animals (*e.g.*, rabbit, goat, mouse or other mammal) may be immunized by one or more injections with the native protein, a synthetic variant thereof, or a derivative of the foregoing. An appropriate immunogenic preparation can contain, for example, the naturally occurring immunogenic protein, a chemically synthesized polypeptide representing the immunogenic protein, or a recombinantly expressed immunogenic protein. Furthermore, the protein may be conjugated to a second protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. The preparation can further include an adjuvant. Various adjuvants used to increase the immunological response include, but are not limited to, Freund's (complete and incomplete), mineral gels (*e.g.*, aluminum hydroxide), surface active substances (*e.g.*, lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, dinitrophenol, etc.), adjuvants usable in humans such as Bacille Calmette-Guerin and *Corynebacterium parvum*, or similar immunostimulatory agents. Additional examples of adjuvants which can be employed include MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate).

The polyclonal antibody molecules directed against the immunogenic protein can be isolated from the mammal (*e.g.*, from the blood) and further purified by well known techniques, such as affinity chromatography using protein A or protein G, which provide primarily the IgG fraction of immune serum. Subsequently, or alternatively, the specific antigen which is the target of the immunoglobulin sought, or an epitope thereof, may be immobilized on a column to purify the immune specific antibody by immunoaffinity chromatography. Purification of immunoglobulins is discussed, for example, by D.

Wilkinson (The Scientist, published by The Scientist, Inc., Philadelphia PA, Vol. 14, No. 8 (April 17, 2000), pp. 25-28).

### Monoclonal Antibodies

5           The term "monoclonal antibody" (MAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs thus  
10       contain an antigen binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

          Monoclonal antibodies can be prepared using hybridoma methods, such as those described by Kohler and Milstein, *Nature*, 256:495 (1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent  
15       to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes can be immunized in vitro.

          The immunizing agent will typically include the protein antigen, a fragment thereof or a fusion protein thereof. Generally, either peripheral blood lymphocytes are used if cells of  
20       human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, MONOCLONAL ANTIBODIES: PRINCIPLES AND PRACTICE, Academic Press, (1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly  
25       myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells can be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas  
30       typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

          Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are sensitive to a

medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Manassas, Virginia.

Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, *J. Immunol.*, 133:3001 (1984); Brodeur et al., MONOCLONAL ANTIBODY PRODUCTION TECHNIQUES AND APPLICATIONS, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the antigen. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, *Anal. Biochem.*, 107:220 (1980). Preferably, antibodies having a high degree of specificity and a high binding affinity for the target antigen are isolated.

After the desired hybridoma cells are identified, the clones can be subcloned by limiting dilution procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells can be grown in vivo as ascites in a mammal.

The monoclonal antibodies secreted by the subclones can be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies can also be made by recombinant DNA methods, such as those described in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA can be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also can be modified, for example, by substituting the coding sequence for human heavy and

light chain constant domains in place of the homologous murine sequences (U.S. Patent No. 4,816,567; Morrison, *Nature* 368, 812-13 (1994)) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted  
5 for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

### Humanized Antibodies

The antibodies directed against the protein antigens of the invention can further  
10 comprise humanized antibodies or human antibodies. These antibodies are suitable for administration to humans without engendering an immune response by the human against the administered immunoglobulin. Humanized forms of antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')<sub>2</sub> or other antigen-binding subsequences of antibodies) that are principally comprised of the  
15 sequence of a human immunoglobulin, and contain minimal sequence derived from a non-human immunoglobulin. Humanization can be performed following the method of Winter and co-workers (Jones et al., *Nature*, 321:522-525 (1986); Riechmann et al., *Nature*, 332:323-327 (1988); Verhoeven et al., *Science*, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. (See  
20 also U.S. Patent No. 5,225,539.) In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies can also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or  
25 substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., 1986; Riechmann et al., 1988; and Presta, *Curr. Opin. Struct. Biol.*, 2:593-596 (1992)).

## Human Antibodies

Fully human antibodies relate to antibody molecules in which essentially the entire sequences of both the light chain and the heavy chain, including the CDRs, arise from human genes. Such antibodies are termed “human antibodies”, or “fully human antibodies” herein. Human monoclonal antibodies can be prepared by the trioma technique; the human B-cell hybridoma technique (see Kozbor, et al., 1983 *Immunol Today* 4: 72) and the EBV hybridoma technique to produce human monoclonal antibodies (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96). Human monoclonal antibodies may be utilized in the practice of the present invention and may be produced by using human hybridomas (see Cote, et al., 1983. *Proc Natl Acad Sci USA* 80: 2026-2030) or by transforming human B-cells with Epstein Barr Virus in vitro (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96).

In addition, human antibodies can also be produced using additional techniques, including phage display libraries (Hoogenboom and Winter, *J. Mol. Biol.*, 227:381 (1991); Marks et al., *J. Mol. Biol.*, 222:581 (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, for example, in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in Marks et al. *Bio/Technology* 10:779-783 (1992); Lonberg et al. *Nature* 368:856-859 (1994); Morrison *Nature* 368:812-13 (1994); Fishwild et al, *Nature Biotechnology* 14:845-51 (1996); Neuberger *Nature Biotechnology* 14:826 (1996); and Lonberg and Huszar *Intern. Rev. Immunol.* 13:65-93 (1995).

Human antibodies may additionally be produced using transgenic nonhuman animals which are modified so as to produce fully human antibodies rather than the animal's endogenous antibodies in response to challenge by an antigen. (See PCT publication WO94/02602). The endogenous genes encoding the heavy and light immunoglobulin chains in the nonhuman host have been incapacitated, and active loci encoding human heavy and light chain immunoglobulins are inserted into the host's genome. The human genes are incorporated, for example, using yeast artificial chromosomes containing the requisite human



DNA segments. An animal which provides all the desired modifications is then obtained as progeny by crossbreeding intermediate transgenic animals containing fewer than the full complement of the modifications. The preferred embodiment of such a nonhuman animal is a mouse, and is termed the Xenomouse<sup>TM</sup> as disclosed in PCT publications WO 96/33735 and WO 96/34096. This animal produces B cells which secrete fully human immunoglobulins. The antibodies can be obtained directly from the animal after immunization with an immunogen of interest, as, for example, a preparation of a polyclonal antibody, or alternatively from immortalized B cells derived from the animal, such as hybridomas producing monoclonal antibodies. Additionally, the genes encoding the immunoglobulins with human variable regions can be recovered and expressed to obtain the antibodies directly, or can be further modified to obtain analogs of antibodies such as, for example, single chain Fv molecules.

An example of a method of producing a nonhuman host, exemplified as a mouse, lacking expression of an endogenous immunoglobulin heavy chain is disclosed in U.S. Patent No. 5,939,598. It can be obtained by a method including deleting the J segment genes from at least one endogenous heavy chain locus in an embryonic stem cell to prevent rearrangement of the locus and to prevent formation of a transcript of a rearranged immunoglobulin heavy chain locus, the deletion being effected by a targeting vector containing a gene encoding a selectable marker; and producing from the embryonic stem cell a transgenic mouse whose somatic and germ cells contain the gene encoding the selectable marker.

A method for producing an antibody of interest, such as a human antibody, is disclosed in U.S. Patent No. 5,916,771. It includes introducing an expression vector that contains a nucleotide sequence encoding a heavy chain into one mammalian host cell in culture, introducing an expression vector containing a nucleotide sequence encoding a light chain into another mammalian host cell, and fusing the two cells to form a hybrid cell. The hybrid cell expresses an antibody containing the heavy chain and the light chain.

In a further improvement on this procedure, a method for identifying a clinically relevant epitope on an immunogen, and a correlative method for selecting an antibody that binds immunospecifically to the relevant epitope with high affinity, are disclosed in PCT publication WO 99/53049.

### **F<sub>ab</sub> Fragments and Single Chain Antibodies**

According to the invention, techniques can be adapted for the production of single-chain antibodies specific to an antigenic protein of the invention (see e.g., U.S. Patent No. 4,946,778). In addition, methods can be adapted for the construction of F<sub>ab</sub> expression  
5 libraries (see e.g., Huse, et al., 1989 Science 246: 1275-1281) to allow rapid and effective identification of monoclonal F<sub>ab</sub> fragments with the desired specificity for a protein or derivatives, fragments, analogs or homologs thereof. Antibody fragments that contain the idiotypes to a protein antigen may be produced by techniques known in the art including, but not limited to: (i) an F<sub>(ab)<sub>2</sub></sub> fragment produced by pepsin digestion of an antibody molecule;  
10 (ii) an F<sub>ab</sub> fragment generated by reducing the disulfide bridges of an F<sub>(ab)<sub>2</sub></sub> fragment; (iii) an F<sub>ab</sub> fragment generated by the treatment of the antibody molecule with papain and a reducing agent and (iv) F<sub>v</sub> fragments

### **Bispecific Antibodies**

Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that  
15 have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for an antigenic protein of the invention. The second binding target is any other antigen, and advantageously is a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the  
20 recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities (Milstein and Cuello, *Nature*, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the  
25 correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker *et al.*, 1991 *EMBO J.*, 10:3655-3659.

Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion  
30 preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the

immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. For further details of generating bispecific antibodies see, for example, Suresh et al., *Methods in Enzymology*, 121:210 (1986).

According to another approach described in WO 96/27011, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers which are recovered from recombinant cell culture. The preferred interface comprises at least a part of the CH3 region of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g. tyrosine or tryptophan). Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g. alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers.

Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g.  $F(ab')_2$  bispecific antibodies). Techniques for generating bispecific antibodies from antibody fragments have been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., *Science* 229:81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate  $F(ab')_2$  fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The  $Fab'$  fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the  $Fab'$ -TNB derivatives is then reconverted to the  $Fab'$ -thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other  $Fab'$ -TNB derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

Additionally,  $Fab'$  fragments can be directly recovered from *E. coli* and chemically coupled to form bispecific antibodies. Shalaby et al., *J. Exp. Med.* 175:217-225 (1992) describe the production of a fully humanized bispecific antibody  $F(ab')_2$  molecule. Each  $Fab'$  fragment was separately secreted from *E. coli* and subjected to directed chemical coupling in vitro to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the ErbB2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets.

Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies

have been produced using leucine zippers. Kostelny et al., *J. Immunol.* 148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers were reduced at the hinge region to form monomers and then re-oxidized to form the antibody

5 heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology described by Hollinger et al., *Proc. Natl. Acad. Sci. USA* 90:6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain ( $V_H$ ) connected to a light-chain variable domain ( $V_L$ ) by a linker which is too short to allow pairing between the

10 two domains on the same chain. Accordingly, the  $V_H$  and  $V_L$  domains of one fragment are forced to pair with the complementary  $V_L$  and  $V_H$  domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See, Gruber et al., *J. Immunol.* 152:5368 (1994).

15 Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al., *J. Immunol.* 147:60 (1991).

Exemplary bispecific antibodies can bind to two different epitopes, at least one of which originates in the protein antigen of the invention. Alternatively, an anti-antigenic arm of an immunoglobulin molecule can be combined with an arm which binds to a triggering

20 molecule on a leukocyte such as a T-cell receptor molecule (e.g. CD2, CD3, CD28, or B7), or Fc receptors for IgG (FcγR), such as FcγRI (CD64), FcγRII (CD32) and FcγRIII (CD16) so as to focus cellular defense mechanisms to the cell expressing the particular antigen.

Bispecific antibodies can also be used to direct cytotoxic agents to cells which express a particular antigen. These antibodies possess an antigen-binding arm and an arm which binds

25 a cytotoxic agent or a radionuclide chelator, such as EOTUBE, DPTA, DOTA, or TETA. Another bispecific antibody of interest binds the protein antigen described herein and further binds tissue factor (TF).

### **Heteroconjugate Antibodies**

30 Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (U.S. Patent No. 4,676,980), and for treatment of HIV infection (WO 91/00360; WO

92/200373; EP 03089). It is contemplated that the antibodies can be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins can be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include  
5 iminothiolate and methyl-4-mercaptobutyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

### **Effector Function Engineering**

It can be desirable to modify the antibody of the invention with respect to effector  
10 function, so as to enhance, e.g., the effectiveness of the antibody in treating cancer. For example, cysteine residue(s) can be introduced into the Fc region, thereby allowing interchain disulfide bond formation in this region. The homodimeric antibody thus generated can have improved internalization capability and/or increased complement-mediated cell killing and antibody-dependent cellular cytotoxicity (ADCC). See Caron et al., *J. Exp Med.*, 176:1191-  
15 1195 (1992) and Shopes, *J. Immunol.*, 148:2918-2922 (1992). Homodimeric antibodies with enhanced anti-tumor activity can also be prepared using heterobifunctional cross-linkers as described in Wolff et al. *Cancer Research*, 53:2560-2565 (1993). Alternatively, an antibody can be engineered that has dual Fc regions and can thereby have enhanced complement lysis and ADCC capabilities. See Stevenson et al., *Anti-Cancer Drug Design*, 3:219-230 (1989).

### **Immunoconjugates**

The invention also pertains to immunoconjugates comprising an antibody conjugated to a cytotoxic agent such as a chemotherapeutic agent, toxin (e.g., an enzymatically active toxin of bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive  
25 isotope (i.e., a radioconjugate).

Chemotherapeutic agents useful in the generation of such immunoconjugates have been described above. Enzymatically active toxins and fragments thereof that can be used include diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from *Pseudomonas aeruginosa*), ricin A chain, abrin A chain, modeccin A chain,  
30 alpha-sarcin, Aleurites fordii proteins, dianthin proteins, Phytolacca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, saponaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes. A

variety of radionuclides are available for the production of radioconjugated antibodies.

Examples include  $^{212}\text{Bi}$ ,  $^{131}\text{I}$ ,  $^{131}\text{In}$ ,  $^{90}\text{Y}$ , and  $^{186}\text{Re}$ .

Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutaraldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as tolyene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta et al., Science, 238:1098 (1987). Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See WO94/11026.

In another embodiment, the antibody can be conjugated to a "receptor" (such as streptavidin) for utilization in tumor pretargeting wherein the antibody-receptor conjugate is administered to the patient, followed by removal of unbound conjugate from the circulation using a clearing agent and then administration of a "ligand" (e.g., avidin) that is in turn conjugated to a cytotoxic agent.

In one embodiment, methods for the screening of antibodies that possess the desired specificity include, but are not limited to, enzyme-linked immunosorbent assay (ELISA) and other immunologically-mediated techniques known within the art. In a specific embodiment, selection of antibodies that are specific to a particular domain of a GPCR protein is facilitated by generation of hybridomas that bind to the fragment of a GPCR protein possessing such a domain. Thus, antibodies that are specific for a desired domain within a GPCR protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

Anti-GPCR antibodies may be used in methods known within the art relating to the localization and/or quantitation of a GPCR protein (e.g., for use in measuring levels of the GPCR protein within appropriate physiological samples, for use in diagnostic methods, for use in imaging the protein, and the like). In a given embodiment, antibodies for GPCR proteins, or derivatives, fragments, analogs or homologs thereof, that contain the antibody derived binding domain, are utilized as pharmacologically-active compounds (hereinafter "Therapeutics").

An anti-GPCR<sub>X</sub> antibody (*e.g.*, monoclonal antibody) can be used to isolate an GPCR<sub>X</sub> polypeptide by standard techniques, such as affinity chromatography or immunoprecipitation. An anti-GPCR<sub>X</sub> antibody can facilitate the purification of natural GPCR<sub>X</sub> polypeptide from cells and of recombinantly-produced GPCR<sub>X</sub> polypeptide expressed in host cells. Moreover, an anti-GPCR<sub>X</sub> antibody can be used to detect GPCR<sub>X</sub> protein (*e.g.*, in a cellular lysate or cell supernatant) in order to evaluate the abundance and pattern of expression of the GPCR<sub>X</sub> protein. Anti-GPCR<sub>X</sub> antibodies can be used diagnostically to monitor protein levels in tissue as part of a clinical testing procedure, *e.g.*, to, for example, determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling (*i.e.*, physically linking) the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, and radioactive materials. Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase,  $\beta$ -galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin, and examples of suitable radioactive material include  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{35}\text{S}$  or  $^3\text{H}$ .

### **GPCR<sub>X</sub> Recombinant Expression Vectors and Host Cells**

Another aspect of the invention pertains to vectors, preferably expression vectors, containing a nucleic acid encoding an GPCR<sub>X</sub> protein, or derivatives, fragments, analogs or homologs thereof. As used herein, the term "vector" refers to a nucleic acid molecule capable of transporting another nucleic acid to which it has been linked. One type of vector is a "plasmid", which refers to a circular double stranded DNA loop into which additional DNA segments can be ligated. Another type of vector is a viral vector, wherein additional DNA segments can be ligated into the viral genome. Certain vectors are capable of autonomous replication in a host cell into which they are introduced (*e.g.*, bacterial vectors having a bacterial origin of replication and episomal mammalian vectors). Other vectors (*e.g.*, non-episomal mammalian vectors) are integrated into the genome of a host cell upon introduction into the host cell, and thereby are replicated along with the host genome. Moreover, certain vectors are capable of directing the expression of genes to which they are

operatively-linked. Such vectors are referred to herein as "expression vectors". In general, expression vectors of utility in recombinant DNA techniques are often in the form of plasmids. In the present specification, "plasmid" and "vector" can be used interchangeably as the plasmid is the most commonly used form of vector. However, the invention is intended to include such other forms of expression vectors, such as viral vectors (*e.g.*, replication defective retroviruses, adenoviruses and adeno-associated viruses), which serve equivalent functions.

The recombinant expression vectors of the invention comprise a nucleic acid of the invention in a form suitable for expression of the nucleic acid in a host cell, which means that the recombinant expression vectors include one or more regulatory sequences, selected on the basis of the host cells to be used for expression, that is operatively-linked to the nucleic acid sequence to be expressed. Within a recombinant expression vector, "operably-linked" is intended to mean that the nucleotide sequence of interest is linked to the regulatory sequence(s) in a manner that allows for expression of the nucleotide sequence (*e.g.*, in an *in vitro* transcription/translation system or in a host cell when the vector is introduced into the host cell).

The term "regulatory sequence" is intended to include promoters, enhancers and other expression control elements (*e.g.*, polyadenylation signals). Such regulatory sequences are described, for example, in Goeddel, GENE EXPRESSION TECHNOLOGY: METHODS IN ENZYMOLOGY 185, Academic Press, San Diego, Calif. (1990). Regulatory sequences include those that direct constitutive expression of a nucleotide sequence in many types of host cell and those that direct expression of the nucleotide sequence only in certain host cells (*e.g.*, tissue-specific regulatory sequences). It will be appreciated by those skilled in the art that the design of the expression vector can depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc. The expression vectors of the invention can be introduced into host cells to thereby produce proteins or peptides, including fusion proteins or peptides, encoded by nucleic acids as described herein (*e.g.*, GPCR proteins, mutant forms of GPCR proteins, fusion proteins, etc.).

The recombinant expression vectors of the invention can be designed for expression of GPCR proteins in prokaryotic or eukaryotic cells. For example, GPCR proteins can be expressed in bacterial cells such as *Escherichia coli*, insect cells (using baculovirus expression vectors) yeast cells or mammalian cells. Suitable host cells are discussed further in Goeddel, GENE EXPRESSION TECHNOLOGY: METHODS IN ENZYMOLOGY 185, Academic Press, San Diego, Calif. (1990). Alternatively, the recombinant expression vector can be



transcribed and translated *in vitro*, for example using T7 promoter regulatory sequences and T7 polymerase.

Expression of proteins in prokaryotes is most often carried out in *Escherichia coli* with vectors containing constitutive or inducible promoters directing the expression of either fusion or non-fusion proteins. Fusion vectors add a number of amino acids to a protein encoded therein, usually to the amino terminus of the recombinant protein. Such fusion vectors typically serve three purposes: (i) to increase expression of recombinant protein; (ii) to increase the solubility of the recombinant protein; and (iii) to aid in the purification of the recombinant protein by acting as a ligand in affinity purification. Often, in fusion expression vectors, a proteolytic cleavage site is introduced at the junction of the fusion moiety and the recombinant protein to enable separation of the recombinant protein from the fusion moiety subsequent to purification of the fusion protein. Such enzymes, and their cognate recognition sequences, include Factor Xa, thrombin and enterokinase. Typical fusion expression vectors include pGEX (Pharmacia Biotech Inc; Smith and Johnson, 1988. *Gene* 67: 31-40), pMAL (New England Biolabs, Beverly, Mass.) and pRIT5 (Pharmacia, Piscataway, N.J.) that fuse glutathione S-transferase (GST), maltose E binding protein, or protein A, respectively, to the target recombinant protein.

Examples of suitable inducible non-fusion *E. coli* expression vectors include pTrc (Amrann *et al.*, (1988) *Gene* 69:301-315) and pET 11d (Studier *et al.*, GENE EXPRESSION TECHNOLOGY: METHODS IN ENZYMOLOGY 185, Academic Press, San Diego, Calif. (1990) 60-89).

One strategy to maximize recombinant protein expression in *E. coli* is to express the protein in a host bacteria with an impaired capacity to proteolytically cleave the recombinant protein. See, e.g., Gottesman, GENE EXPRESSION TECHNOLOGY: METHODS IN ENZYMOLOGY 185, Academic Press, San Diego, Calif. (1990) 119-128. Another strategy is to alter the nucleic acid sequence of the nucleic acid to be inserted into an expression vector so that the individual codons for each amino acid are those preferentially utilized in *E. coli* (see, e.g., Wada, *et al.*, 1992 *Nucl. Acids Res.* 20:2111-2118). Such alteration of nucleic acid sequences of the invention can be carried out by standard DNA synthesis techniques.

In another embodiment, the GPCR expression vector is a yeast expression vector. Examples of vectors for expression in yeast *Saccharomyces cerevisiae* include pYepSec1 (Baldari, *et al.*, 1987 *EMBO J.* 6: 229-234), pMFa (Kurjan and Herskowitz, 1982 *Cell* 30:933-943), pJRY88 (Schultz *et al.*, 1987 *Gene* 54:113-123), pYES2 (Invitrogen Corporation, San Diego, Calif.), and picZ (Invitrogen Corp, San Diego, Calif.).

Alternatively, GPCR<sub>X</sub> can be expressed in insect cells using baculovirus expression vectors. Baculovirus vectors available for expression of proteins in cultured insect cells (*e.g.*, SF9 cells) include the pAc series (Smith, *et al.*, 1983 *Mol. Cell. Biol.* 3:2156-2165) and the pVL series (Lucklow and Summers, 1989 *Virology* 170:31-39).

5           In yet another embodiment, a nucleic acid of the invention is expressed in mammalian cells using a mammalian expression vector. Examples of mammalian expression vectors include pCDM8 (Seed, 1987 *Nature* 329:840) and pMT2PC (Kaufman, *et al.*, 1987 *EMBO J.* 6:187-195). When used in mammalian cells, the expression vector's control functions are often provided by viral regulatory elements. For example, commonly used  
10           promoters are derived from polyoma, adenovirus 2, cytomegalovirus, and simian virus 40. For other suitable expression systems for both prokaryotic and eukaryotic cells see, *e.g.*, Chapters 16 and 17 of Sambrook, *et al.*, MOLECULAR CLONING: A LABORATORY MANUAL. 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1989.

15           In another embodiment, the recombinant mammalian expression vector is capable of directing expression of the nucleic acid preferentially in a particular cell type (*e.g.*, tissue-specific regulatory elements are used to express the nucleic acid). Tissue-specific regulatory elements are known in the art. Non-limiting examples of suitable tissue-specific promoters include the albumin promoter (liver-specific; Pinkert, *et al.*, 1987 *Genes Dev.*  
20           1:268-277), lymphoid-specific promoters (Calame and Eaton, 1988 *Adv. Immunol.* 43:235-275), in particular promoters of T cell receptors (Winoto and Baltimore, 1989 *EMBO J.* 8:729-733) and immunoglobulins (Banerji, *et al.*, 1983 *Cell* 33:729-740; Queen and Baltimore, 1983 *Cell* 33:741-748), neuron-specific promoters (*e.g.*, the neurofilament promoter; Byrne and Ruddle, 1989 *Proc. Natl. Acad. Sci. USA* 86:5473-5477),  
25           pancreas-specific promoters (Edlund, *et al.*, 1985 *Science* 230:912-916), and mammary gland-specific promoters (*e.g.*, milk whey promoter; U.S. Pat. No. 4,873,316 and European Application Publication No. 264,166). Developmentally-regulated promoters are also encompassed, *e.g.*, the murine hox promoters (Kessel and Gruss, 1990 *Science* 249:374-379) and the  $\alpha$ -fetoprotein promoter (Campes and Tilghman, 1989 *Genes Dev.* 3:537-546).

30           The invention further provides a recombinant expression vector comprising a DNA molecule of the invention cloned into the expression vector in an antisense orientation. That is, the DNA molecule is operatively-linked to a regulatory sequence in a manner that allows for expression (by transcription of the DNA molecule) of an RNA molecule that is antisense to GPCR<sub>X</sub> mRNA. Regulatory sequences operatively linked to a nucleic acid cloned in the

antisense orientation can be chosen that direct the continuous expression of the antisense RNA molecule in a variety of cell types, for instance viral promoters and/or enhancers, or regulatory sequences can be chosen that direct constitutive, tissue specific or cell type specific expression of antisense RNA. The antisense expression vector can be in the form of a recombinant plasmid, phagemid or attenuated virus in which antisense nucleic acids are produced under the control of a high efficiency regulatory region, the activity of which can be determined by the cell type into which the vector is introduced. For a discussion of the regulation of gene expression using antisense genes *see, e.g., Weintraub, et al., "Antisense RNA as a molecular tool for genetic analysis," Reviews-Trends in Genetics, Vol. 1(1) 1986.*

Another aspect of the invention pertains to host cells into which a recombinant expression vector of the invention has been introduced. The terms "host cell" and "recombinant host cell" are used interchangeably herein. It is understood that such terms refer not only to the particular subject cell but also to the progeny or potential progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term as used herein.

A host cell can be any prokaryotic or eukaryotic cell. For example, GPCR<sub>X</sub> protein can be expressed in bacterial cells such as *E. coli*, insect cells, yeast or mammalian cells (such as Chinese hamster ovary cells (CHO) or COS cells). Other suitable host cells are known to those skilled in the art.

Vector DNA can be introduced into prokaryotic or eukaryotic cells via conventional transformation or transfection techniques. As used herein, the terms "transformation" and "transfection" are intended to refer to a variety of art-recognized techniques for introducing foreign nucleic acid (*e.g., DNA*) into a host cell, including calcium phosphate or calcium chloride co-precipitation, DEAE-dextran-mediated transfection, lipofection, or electroporation. Suitable methods for transforming or transfecting host cells can be found in Sambrook, *et al.* (MOLECULAR CLONING: A LABORATORY MANUAL. 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1989), and other laboratory manuals.

For stable transfection of mammalian cells, it is known that, depending upon the expression vector and transfection technique used, only a small fraction of cells may integrate the foreign DNA into their genome. In order to identify and select these integrants, a gene that encodes a selectable marker (*e.g., resistance to antibiotics*) is generally introduced into the host cells along with the gene of interest. Various selectable markers include those that

confer resistance to drugs, such as G418, hygromycin and methotrexate. Nucleic acid encoding a selectable marker can be introduced into a host cell on the same vector as that encoding GPCR<sub>X</sub> or can be introduced on a separate vector. Cells stably transfected with the introduced nucleic acid can be identified by drug selection (*e.g.*, cells that have incorporated the selectable marker gene will survive, while the other cells die).

A host cell of the invention, such as a prokaryotic or eukaryotic host cell in culture, can be used to produce (*i.e.*, express) GPCR<sub>X</sub> protein. Accordingly, the invention further provides methods for producing GPCR<sub>X</sub> protein using the host cells of the invention. In one embodiment, the method comprises culturing the host cell of invention (into which a recombinant expression vector encoding GPCR<sub>X</sub> protein has been introduced) in a suitable medium such that GPCR<sub>X</sub> protein is produced. In another embodiment, the method further comprises isolating GPCR<sub>X</sub> protein from the medium or the host cell.

### **Transgenic GPCR<sub>X</sub> Animals**

The host cells of the invention can also be used to produce non-human transgenic animals. For example, in one embodiment, a host cell of the invention is a fertilized oocyte or an embryonic stem cell into which GPCR<sub>X</sub> protein-coding sequences have been introduced. Such host cells can then be used to create non-human transgenic animals in which exogenous GPCR<sub>X</sub> sequences have been introduced into their genome or homologous recombinant animals in which endogenous GPCR<sub>X</sub> sequences have been altered. Such animals are useful for studying the function and/or activity of GPCR<sub>X</sub> protein and for identifying and/or evaluating modulators of GPCR<sub>X</sub> protein activity. As used herein, a "transgenic animal" is a non-human animal, preferably a mammal, more preferably a rodent such as a rat or mouse, in which one or more of the cells of the animal includes a transgene. Other examples of transgenic animals include non-human primates, sheep, dogs, cows, goats, chickens, amphibians, etc. A transgene is exogenous DNA that is integrated into the genome of a cell from which a transgenic animal develops and that remains in the genome of the mature animal, thereby directing the expression of an encoded gene product in one or more cell types or tissues of the transgenic animal. As used herein, a "homologous recombinant animal" is a non-human animal, preferably a mammal, more preferably a mouse, in which an endogenous GPCR<sub>X</sub> gene has been altered by homologous recombination between the endogenous gene and an exogenous DNA molecule introduced into a cell of the animal, *e.g.*, an embryonic cell of the animal, prior to development of the animal.

A transgenic animal of the invention can be created by introducing GPCR<sub>X</sub>-encoding nucleic acid into the male pronuclei of a fertilized oocyte (*e.g.*, by microinjection, retroviral infection) and allowing the oocyte to develop in a pseudopregnant female foster animal. The human GPCR<sub>X</sub> cDNA sequences of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 can be introduced as a transgene into the genome of a non-human animal. Alternatively, a non-human homologue of the human GPCR<sub>X</sub> gene, such as a mouse GPCR<sub>X</sub> gene, can be isolated based on hybridization to the human GPCR<sub>X</sub> cDNA (described further *supra*) and used as a transgene. Intronic sequences and polyadenylation signals can also be included in the transgene to increase the efficiency of expression of the transgene. A tissue-specific regulatory sequence(s) can be operably-linked to the GPCR<sub>X</sub> transgene to direct expression of GPCR<sub>X</sub> protein to particular cells. Methods for generating transgenic animals via embryo manipulation and microinjection, particularly animals such as mice, have become conventional in the art and are described, for example, in U.S. Patent Nos. 4,736,866; 4,870,009; and 4,873,191; and Hogan, 1986. In: MANIPULATING THE MOUSE EMBRYO, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. Similar methods are used for production of other transgenic animals. A transgenic founder animal can be identified based upon the presence of the GPCR<sub>X</sub> transgene in its genome and/or expression of GPCR<sub>X</sub> mRNA in tissues or cells of the animals. A transgenic founder animal can then be used to breed additional animals carrying the transgene. Moreover, transgenic animals carrying a transgene-encoding GPCR<sub>X</sub> protein can further be bred to other transgenic animals carrying other transgenes.

To create a homologous recombinant animal, a vector is prepared which contains at least a portion of an GPCR<sub>X</sub> gene into which a deletion, addition or substitution has been introduced to thereby alter, *e.g.*, functionally disrupt, the GPCR<sub>X</sub> gene. The GPCR<sub>X</sub> gene can be a human gene (*e.g.*, the cDNA of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39), but more preferably, is a non-human homologue of a human GPCR<sub>X</sub> gene. For example, a mouse homologue of human GPCR<sub>X</sub> gene of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 can be used to construct a homologous recombination vector suitable for altering an endogenous GPCR<sub>X</sub> gene in the mouse genome. In one embodiment, the vector is designed such that, upon homologous recombination, the endogenous GPCR<sub>X</sub> gene is functionally disrupted (*i.e.*, no longer encodes a functional protein; also referred to as a "knock out" vector).

Alternatively, the vector can be designed such that, upon homologous recombination, the endogenous GPCR<sub>X</sub> gene is mutated or otherwise altered but still encodes functional

protein (*e.g.*, the upstream regulatory region can be altered to thereby alter the expression of the endogenous GPCR<sub>X</sub> protein). In the homologous recombination vector, the altered portion of the GPCR<sub>X</sub> gene is flanked at its 5'- and 3'-termini by additional nucleic acid of the GPCR<sub>X</sub> gene to allow for homologous recombination to occur between the exogenous

5 GPCR<sub>X</sub> gene carried by the vector and an endogenous GPCR<sub>X</sub> gene in an embryonic stem cell. The additional flanking GPCR<sub>X</sub> nucleic acid is of sufficient length for successful homologous recombination with the endogenous gene. Typically, several kilobases of flanking DNA (both at the 5'- and 3'-termini) are included in the vector. *See, e.g.*, Thomas, *et al.*, 1987. *Cell* 51: 503 for a description of homologous recombination vectors. The vector is

10 ten introduced into an embryonic stem cell line (*e.g.*, by electroporation) and cells in which the introduced GPCR<sub>X</sub> gene has homologously-recombined with the endogenous GPCR<sub>X</sub> gene are selected. *See, e.g.*, Li, *et al.*, 1992 *Cell* 69:915.

The selected cells are then injected into a blastocyst of an animal (*e.g.*, a mouse) to form aggregation chimeras. *See, e.g.*, Bradley, 1987. In: TERATOCARCINOMAS AND

15 EMBRYONIC STEM CELLS: A PRACTICAL APPROACH, Robertson, ed. IRL, Oxford, pp. 113-152. A chimeric embryo can then be implanted into a suitable pseudopregnant female foster animal and the embryo brought to term. Progeny harboring the homologously-recombined DNA in their germ cells can be used to breed animals in which all cells of the animal contain the homologously-recombined DNA by germline transmission of the

20 transgene. Methods for constructing homologous recombination vectors and homologous recombinant animals are described further in Bradley, 1991. *Curr. Opin. Biotechnol.* 2: 823-829; PCT International Publication Nos.: WO 90/11354; WO 91/01140; WO 92/0968; and WO 93/04169.

In another embodiment, transgenic non-humans animals can be produced that contain

25 selected systems that allow for regulated expression of the transgene. One example of such a system is the cre/loxP recombinase system of bacteriophage P1. For a description of the cre/loxP recombinase system, *See, e.g.*, Lakso, *et al.*, 1992 *Proc. Natl. Acad. Sci. USA* 89:6232-6236. Another example of a recombinase system is the FLP recombinase system of *Saccharomyces cerevisiae*. *See, O'Gorman, et al.*, 1991 *Science* 251:1351-1355. If a

30 cre/loxP recombinase system is used to regulate expression of the transgene, animals containing transgenes encoding both the Cre recombinase and a selected protein are required. Such animals can be provided through the construction of "double" transgenic animals, *e.g.*, by mating two transgenic animals, one containing a transgene encoding a selected protein and the other containing a transgene encoding a recombinase.

Clones of the non-human transgenic animals described herein can also be produced according to the methods described in Wilmut, *et al.*, 1997 *Nature* 385:810-813. In brief, a cell (*e.g.*, a somatic cell) from the transgenic animal can be isolated and induced to exit the growth cycle and enter G<sub>0</sub> phase. The quiescent cell can then be fused, *e.g.*, through the use of electrical pulses, to an enucleated oocyte from an animal of the same species from which the quiescent cell is isolated. The reconstructed oocyte is then cultured such that it develops to morula or blastocyte and then transferred to pseudopregnant female foster animal. The offspring borne of this female foster animal will be a clone of the animal from which the cell (*e.g.*, the somatic cell) is isolated.

### Pharmaceutical Compositions

The GPCR<sub>X</sub> nucleic acid molecules, GPCR<sub>X</sub> proteins, and anti-GPCR<sub>X</sub> antibodies (also referred to herein as "active compounds") of the invention, and derivatives, fragments, analogs and homologs thereof, can be incorporated into pharmaceutical compositions suitable for administration. Such compositions typically comprise the nucleic acid molecule, protein, or antibody and a pharmaceutically acceptable carrier. As used herein, "pharmaceutically acceptable carrier" is intended to include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like, compatible with pharmaceutical administration. Suitable carriers are described in the most recent edition of Remington's Pharmaceutical Sciences, a standard reference text in the field, which is incorporated herein by reference. Preferred examples of such carriers or diluents include, but are not limited to, water, saline, finger's solutions, dextrose solution, and 5% human serum albumin. Liposomes and non-aqueous vehicles such as fixed oils may also be used. The use of such media and agents for pharmaceutically active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active compound, use thereof in the compositions is contemplated. Supplementary active compounds can also be incorporated into the compositions.

A pharmaceutical composition of the invention is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, *e.g.*, intravenous, intradermal, subcutaneous, oral (*e.g.*, inhalation), transdermal (*i.e.*, topical), transmucosal, and rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or

methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid (EDTA); buffers such as acetates, citrates or phosphates, and agents for the adjustment of tonicity such as sodium chloride or dextrose. The pH can be adjusted with acids or bases, such as hydrochloric acid or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL™ (BASF, Parsippany, N.J.) or phosphate buffered saline (PBS). In all cases, the composition must be sterile and should be fluid to the extent that easy syringeability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, polyalcohols such as manitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent which delays absorption, for example, aluminum monostearate and gelatin.

Sterile injectable solutions can be prepared by incorporating the active compound (e.g., an GPCR<sub>X</sub> protein or anti-GPCR<sub>X</sub> antibody) in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, methods of preparation are vacuum drying and freeze-drying that yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.



Oral compositions generally include an inert diluent or an edible carrier. They can be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules. Oral compositions can also be prepared using a fluid carrier for use as a mouthwash, wherein the compound in the fluid carrier is applied orally and swished and expectorated or swallowed. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition. The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring.

For administration by inhalation, the compounds are delivered in the form of an aerosol spray from pressured container or dispenser which contains a suitable propellant, *e.g.*, a gas such as carbon dioxide, or a nebulizer.

Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in the art.

The compounds can also be prepared in the form of suppositories (*e.g.*, with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

In one embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to

viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811.

5 It is especially advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the invention are dictated by and directly  
10 dependent on the unique characteristics of the active compound and the particular therapeutic effect to be achieved, and the limitations inherent in the art of compounding such an active compound for the treatment of individuals.

The nucleic acid molecules of the invention can be inserted into vectors and used as gene therapy vectors. Gene therapy vectors can be delivered to a subject by, for example,  
15 intravenous injection, local administration (*see, e.g.*, U.S. Patent No. 5,328,470) or by stereotactic injection (*see, e.g.*, Chen, *et al.*, 1994 *Proc. Natl. Acad. Sci. USA* 91:3054-3057). The pharmaceutical preparation of the gene therapy vector can include the gene therapy vector in an acceptable diluent, or can comprise a slow release matrix in which the gene delivery vehicle is imbedded. Alternatively, where the complete gene delivery vector can be  
20 produced intact from recombinant cells, *e.g.*, retroviral vectors, the pharmaceutical preparation can include one or more cells that produce the gene delivery system.

The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

### Screening and Detection Methods

25 The isolated nucleic acid molecules of the invention can be used to express GPCR<sub>X</sub> protein (*e.g.*, via a recombinant expression vector in a host cell in gene therapy applications), to detect GPCR<sub>X</sub> mRNA (*e.g.*, in a biological sample) or a genetic lesion in an GPCR<sub>X</sub> gene, and to modulate GPCR<sub>X</sub> activity, as described further, below. In addition, the GPCR<sub>X</sub> proteins can be used to screen drugs or compounds that modulate the GPCR<sub>X</sub> protein activity  
30 or expression as well as to treat disorders characterized by insufficient or excessive production of GPCR<sub>X</sub> protein or production of GPCR<sub>X</sub> protein forms that have decreased or aberrant activity compared to GPCR<sub>X</sub> wild-type protein (*e.g.*; diabetes (regulates insulin release); obesity (binds and transport lipids); metabolic disturbances associated with obesity,

the metabolic syndrome X, as well as anorexia and wasting disorders associated with chronic diseases and various cancers, and infectious disease (possesses anti-microbial activity) and the various dyslipidemias. In addition, the anti-GPCR<sub>X</sub> antibodies of the invention can be used to detect and isolate GPCR<sub>X</sub> proteins and modulate GPCR<sub>X</sub> activity. In yet a further  
5 aspect, the invention can be used in methods to influence appetite, absorption of nutrients and the disposition of metabolic substrates in both a positive and negative fashion.

The invention further pertains to novel agents identified by the screening assays described herein and uses thereof for treatments as described, *supra*.

### Screening Assays

10 The invention provides a method (also referred to herein as a "screening assay") for identifying modulators, *i.e.*, candidate or test compounds or agents (*e.g.*, peptides, peptidomimetics, small molecules or other drugs) that bind to GPCR<sub>X</sub> proteins or have a stimulatory or inhibitory effect on, *e.g.*, GPCR<sub>X</sub> protein expression or GPCR<sub>X</sub> protein activity. The invention also includes compounds identified in the screening assays described  
15 herein.

In one embodiment, the invention provides assays for screening candidate or test compounds which bind to or modulate the activity of the membrane-bound form of an GPCR<sub>X</sub> protein or polypeptide or biologically-active portion thereof. The test compounds of the invention can be obtained using any of the numerous approaches in combinatorial library  
20 methods known in the art, including: biological libraries; spatially addressable parallel solid phase or solution phase libraries; synthetic library methods requiring deconvolution; the "one-bead one-compound" library method; and synthetic library methods using affinity chromatography selection. The biological library approach is limited to peptide libraries, while the other four approaches are applicable to peptide, non-peptide oligomer or small  
25 molecule libraries of compounds. *See, e.g.*, Lam, 1997 *Anticancer Drug Design* 12:145.

A "small molecule" as used herein, is meant to refer to a composition that has a molecular weight of less than about 5 kD and most preferably less than about 4 kD. Small molecules can be, *e.g.*, nucleic acids, peptides, polypeptides, peptidomimetics, carbohydrates, lipids or other organic or inorganic molecules. Libraries of chemical and/or biological  
30 mixtures, such as fungal, bacterial, or algal extracts, are known in the art and can be screened with any of the assays of the invention.

Examples of methods for the synthesis of molecular libraries can be found in the art, for example in: DeWitt, *et al.*, 1993 *Proc. Natl. Acad. Sci. U.S.A.* 90:6909; Erb, *et al.*, 1994

*Proc. Natl. Acad. Sci. U.S.A.* 91:11422; Zuckermann, *et al.*, 1994 *J. Med. Chem.* 37:2678; Cho, *et al.*, 1993 *Science* 261:1303; Carrell, *et al.*, 1994 *Angew. Chem. Int. Ed. Engl.* 33:2059; Carell, *et al.*, 1994 *Angew. Chem. Int. Ed. Engl.* 33:2061; and Gallop, *et al.*, 1994 *J. Med. Chem.* 37:1233.

5 Libraries of compounds may be presented in solution (*e.g.*, Houghten, 1992 *Biotechniques* 13:412-421), or on beads (Lam, 1991 *Nature* 354:82-84), on chips (Fodor, 1993 *Nature* 364:555-556), bacteria (Ladner, U.S. Patent No. 5,223,409), spores (Ladner, U.S. Patent 5,233,409), plasmids (Cull, *et al.*, 1992 *Proc. Natl. Acad. Sci. USA* 89:1865-1869) or on phage (Scott and Smith, 1990. *Science* 249: 386-390; Devlin, 1990  
10 *Science* 249:404-406; Cwirla, *et al.*, 1990 *Proc. Natl. Acad. Sci. U.S.A.* 87:6378-6382; Felici, 1991 *J. Mol. Biol.* 222:301-310; Ladner, U.S. Patent No. 5,233,409.).

In one embodiment, an assay is a cell-based assay in which a cell which expresses a membrane-bound form of GPCR<sub>X</sub> protein, or a biologically-active portion thereof, on the cell surface is contacted with a test compound and the ability of the test compound to bind to  
15 an GPCR<sub>X</sub> protein determined. The cell, for example, can of mammalian origin or a yeast cell. Determining the ability of the test compound to bind to the GPCR<sub>X</sub> protein can be accomplished, for example, by coupling the test compound with a radioisotope or enzymatic label such that binding of the test compound to the GPCR<sub>X</sub> protein or biologically-active portion thereof can be determined by detecting the labeled compound in a complex. For  
20 example, test compounds can be labeled with <sup>125</sup>I, <sup>35</sup>S, <sup>14</sup>C, or <sup>3</sup>H, either directly or indirectly, and the radioisotope detected by direct counting of radioemission or by scintillation counting. Alternatively, test compounds can be enzymatically-labeled with, for example, horseradish peroxidase, alkaline phosphatase, or luciferase, and the enzymatic label detected by  
25 determination of conversion of an appropriate substrate to product. In one embodiment, the assay comprises contacting a cell which expresses a membrane-bound form of GPCR<sub>X</sub> protein, or a biologically-active portion thereof, on the cell surface with a known compound which binds GPCR<sub>X</sub> to form an assay mixture, contacting the assay mixture with a test compound, and determining the ability of the test compound to interact with an GPCR<sub>X</sub> protein, wherein determining the ability of the test compound to interact with an GPCR<sub>X</sub>  
30 protein comprises determining the ability of the test compound to preferentially bind to GPCR<sub>X</sub> protein or a biologically-active portion thereof as compared to the known compound.

In another embodiment, an assay is a cell-based assay comprising contacting a cell expressing a membrane-bound form of GPCR<sub>X</sub> protein, or a biologically-active portion

thereof, on the cell surface with a test compound and determining the ability of the test compound to modulate (*e.g.*, stimulate or inhibit) the activity of the GPCR<sub>X</sub> protein or biologically-active portion thereof. Determining the ability of the test compound to modulate the activity of GPCR<sub>X</sub> or a biologically-active portion thereof can be accomplished, for  
5 example, by determining the ability of the GPCR<sub>X</sub> protein to bind to or interact with an GPCR<sub>X</sub> target molecule. As used herein, a "target molecule" is a molecule with which an GPCR<sub>X</sub> protein binds or interacts in nature, for example, a molecule on the surface of a cell which expresses an GPCR<sub>X</sub> interacting protein, a molecule on the surface of a second cell, a molecule in the extracellular milieu, a molecule associated with the internal surface of a cell  
10 membrane or a cytoplasmic molecule. An GPCR<sub>X</sub> target molecule can be a non-GPCR<sub>X</sub> molecule or an GPCR<sub>X</sub> protein or polypeptide of the invention. In one embodiment, an GPCR<sub>X</sub> target molecule is a component of a signal transduction pathway that facilitates transduction of an extracellular signal (*e.g.* a signal generated by binding of a compound to a membrane-bound GPCR<sub>X</sub> molecule) through the cell membrane and into the cell. The  
15 target, for example, can be a second intercellular protein that has catalytic activity or a protein that facilitates the association of downstream signaling molecules with GPCR<sub>X</sub>.

Determining the ability of the GPCR<sub>X</sub> protein to bind to or interact with an GPCR<sub>X</sub> target molecule can be accomplished by one of the methods described above for determining direct binding. In one embodiment, determining the ability of the GPCR<sub>X</sub> protein to bind to  
20 or interact with an GPCR<sub>X</sub> target molecule can be accomplished by determining the activity of the target molecule. For example, the activity of the target molecule can be determined by detecting induction of a cellular second messenger of the target (*i.e.* intracellular Ca<sup>2+</sup>, diacylglycerol, IP<sub>3</sub>, etc.), detecting catalytic/enzymatic activity of the target an appropriate substrate, detecting the induction of a reporter gene (comprising an GPCR<sub>X</sub>-responsive  
25 regulatory element operatively linked to a nucleic acid encoding a detectable marker, *e.g.*, luciferase), or detecting a cellular response, for example, cell survival, cellular differentiation, or cell proliferation.

In yet another embodiment, an assay of the invention is a cell-free assay comprising contacting an GPCR<sub>X</sub> protein or biologically-active portion thereof with a test compound and  
30 determining the ability of the test compound to bind to the GPCR<sub>X</sub> protein or biologically-active portion thereof. Binding of the test compound to the GPCR<sub>X</sub> protein can be determined either directly or indirectly as described above. In one such embodiment, the assay comprises contacting the GPCR<sub>X</sub> protein or biologically-active portion thereof with a known compound which binds GPCR<sub>X</sub> to form an assay mixture, contacting the assay

mixture with a test compound, and determining the ability of the test compound to interact with an GPCR<sub>X</sub> protein, wherein determining the ability of the test compound to interact with an GPCR<sub>X</sub> protein comprises determining the ability of the test compound to preferentially bind to GPCR<sub>X</sub> or biologically-active portion thereof as compared to the known compound.

In still another embodiment, an assay is a cell-free assay comprising contacting GPCR<sub>X</sub> protein or biologically-active portion thereof with a test compound and determining the ability of the test compound to modulate (*e.g.* stimulate or inhibit) the activity of the GPCR<sub>X</sub> protein or biologically-active portion thereof. Determining the ability of the test compound to modulate the activity of GPCR<sub>X</sub> can be accomplished, for example, by determining the ability of the GPCR<sub>X</sub> protein to bind to an GPCR<sub>X</sub> target molecule by one of the methods described above for determining direct binding. In an alternative embodiment, determining the ability of the test compound to modulate the activity of GPCR<sub>X</sub> protein can be accomplished by determining the ability of the GPCR<sub>X</sub> protein further modulate an GPCR<sub>X</sub> target molecule. For example, the catalytic/enzymatic activity of the target molecule on an appropriate substrate can be determined as described, *supra*.

In yet another embodiment, the cell-free assay comprises contacting the GPCR<sub>X</sub> protein or biologically-active portion thereof with a known compound which binds GPCR<sub>X</sub> protein to form an assay mixture, contacting the assay mixture with a test compound, and determining the ability of the test compound to interact with an GPCR<sub>X</sub> protein, wherein determining the ability of the test compound to interact with an GPCR<sub>X</sub> protein comprises determining the ability of the GPCR<sub>X</sub> protein to preferentially bind to or modulate the activity of an GPCR<sub>X</sub> target molecule.

The cell-free assays of the invention are amenable to use of both the soluble form or the membrane-bound form of GPCR<sub>X</sub> protein. In the case of cell-free assays comprising the membrane-bound form of GPCR<sub>X</sub> protein, it may be desirable to utilize a solubilizing agent such that the membrane-bound form of GPCR<sub>X</sub> protein is maintained in solution. Examples of such solubilizing agents include non-ionic detergents such as n-octylglucoside, n-dodecylglucoside, n-dodecylmaltoside, octanoyl-N-methylglucamide, decanoyl-N-methylglucamide, Triton<sup>®</sup> X-100, Triton<sup>®</sup> X-114, Thesit<sup>®</sup>, Isotridecypoly(ethylene glycol ether)<sub>n</sub>, N-dodecyl--N,N-dimethyl-3-ammonio-1-propane sulfonate, 3-(3-cholamidopropyl) dimethylamminiol-1-propane sulfonate (CHAPS), or 3-(3-cholamidopropyl)dimethylamminiol-2-hydroxy-1-propane sulfonate (CHAPSO).

In more than one embodiment of the above assay methods of the invention, it may be desirable to immobilize either GPCR<sub>X</sub> protein or its target molecule to facilitate separation of complexed from uncomplexed forms of one or both of the proteins, as well as to accommodate automation of the assay. Binding of a test compound to GPCR<sub>X</sub> protein, or interaction of GPCR<sub>X</sub> protein with a target molecule in the presence and absence of a candidate compound, can be accomplished in any vessel suitable for containing the reactants. Examples of such vessels include microtiter plates, test tubes, and micro-centrifuge tubes. In one embodiment, a fusion protein can be provided that adds a domain that allows one or both of the proteins to be bound to a matrix. For example, GST-GPCR<sub>X</sub> fusion proteins or GST-target fusion proteins can be adsorbed onto glutathione sepharose beads (Sigma Chemical, St. Louis, MO) or glutathione derivatized microtiter plates, that are then combined with the test compound or the test compound and either the non-adsorbed target protein or GPCR<sub>X</sub> protein, and the mixture is incubated under conditions conducive to complex formation (*e.g.*, at physiological conditions for salt and pH). Following incubation, the beads or microtiter plate wells are washed to remove any unbound components, the matrix immobilized in the case of beads, complex determined either directly or indirectly, for example, as described, *supra*. Alternatively, the complexes can be dissociated from the matrix, and the level of GPCR<sub>X</sub> protein binding or activity determined using standard techniques.

Other techniques for immobilizing proteins on matrices can also be used in the screening assays of the invention. For example, either the GPCR<sub>X</sub> protein or its target molecule can be immobilized utilizing conjugation of biotin and streptavidin. Biotinylated GPCR<sub>X</sub> protein or target molecules can be prepared from biotin-NHS (N-hydroxy-succinimide) using techniques well-known within the art (*e.g.*, biotinylation kit, Pierce Chemicals, Rockford, Ill.), and immobilized in the wells of streptavidin-coated 96 well plates (Pierce Chemical). Alternatively, antibodies reactive with GPCR<sub>X</sub> protein or target molecules, but which do not interfere with binding of the GPCR<sub>X</sub> protein to its target molecule, can be derivatized to the wells of the plate, and unbound target or GPCR<sub>X</sub> protein trapped in the wells by antibody conjugation. Methods for detecting such complexes, in addition to those described above for the GST-immobilized complexes, include immunodetection of complexes using antibodies reactive with the GPCR<sub>X</sub> protein or target molecule, as well as enzyme-linked assays that rely on detecting an enzymatic activity associated with the GPCR<sub>X</sub> protein or target molecule.

In another embodiment, modulators of GPCR<sub>X</sub> protein expression are identified in a method wherein a cell is contacted with a candidate compound and the expression of GPCR<sub>X</sub>

mRNA or protein in the cell is determined. The level of expression of GPCR<sub>X</sub> mRNA or protein in the presence of the candidate compound is compared to the level of expression of GPCR<sub>X</sub> mRNA or protein in the absence of the candidate compound. The candidate compound can then be identified as a modulator of GPCR<sub>X</sub> mRNA or protein expression based upon this comparison. For example, when expression of GPCR<sub>X</sub> mRNA or protein is greater (*i.e.*, statistically significantly greater) in the presence of the candidate compound than in its absence, the candidate compound is identified as a stimulator of GPCR<sub>X</sub> mRNA or protein expression. Alternatively, when expression of GPCR<sub>X</sub> mRNA or protein is less (statistically significantly less) in the presence of the candidate compound than in its absence, the candidate compound is identified as an inhibitor of GPCR<sub>X</sub> mRNA or protein expression. The level of GPCR<sub>X</sub> mRNA or protein expression in the cells can be determined by methods described herein for detecting GPCR<sub>X</sub> mRNA or protein.

In yet another aspect of the invention, the GPCR<sub>X</sub> proteins can be used as "bait proteins" in a two-hybrid assay or three hybrid assay (*see, e.g.*, U.S. Patent No. 5,283,317; Zervos, *et al.*, 1993 *Cell* 72:223-232; Madura, *et al.*, 1993 *J. Biol. Chem.* 268:12046-12054; Bartel, *et al.*, 1993 *Biotechniques* 14:920-924; Iwabuchi, *et al.*, 1993 *Oncogene* 8:1693-1696; and Brent WO 94/10300), to identify other proteins that bind to or interact with GPCR<sub>X</sub> ("GPCR<sub>X</sub>-binding proteins" or "GPCR<sub>X</sub>-bp") and modulate GPCR<sub>X</sub> activity. Such GPCR<sub>X</sub>-binding proteins are also likely to be involved in the propagation of signals by the GPCR<sub>X</sub> proteins as, for example, upstream or downstream elements of the GPCR<sub>X</sub> pathway.

The two-hybrid system is based on the modular nature of most transcription factors, which consist of separable DNA-binding and activation domains. Briefly, the assay utilizes two different DNA constructs. In one construct, the gene that codes for GPCR<sub>X</sub> is fused to a gene encoding the DNA binding domain of a known transcription factor (*e.g.*, GAL-4). In the other construct, a DNA sequence, from a library of DNA sequences, that encodes an unidentified protein ("prey" or "sample") is fused to a gene that codes for the activation domain of the known transcription factor. If the "bait" and the "prey" proteins are able to interact, *in vivo*, forming an GPCR<sub>X</sub>-dependent complex, the DNA-binding and activation domains of the transcription factor are brought into close proximity. This proximity allows transcription of a reporter gene (*e.g.*, LacZ) that is operably linked to a transcriptional regulatory site responsive to the transcription factor. Expression of the reporter gene can be detected and cell colonies containing the functional transcription factor can be isolated and used to obtain the cloned gene that encodes the protein which interacts with GPCR<sub>X</sub>.



The invention further pertains to novel agents identified by the aforementioned screening assays and uses thereof for treatments as described herein.

### Detection Assays

5            Portions or fragments of the cDNA sequences identified herein (and the corresponding complete gene sequences) can be used in numerous ways as polynucleotide reagents. By way of example, and not of limitation, these sequences can be used to: (i) map their respective genes on a chromosome; and, thus, locate gene regions associated with genetic disease; (ii) identify an individual from a minute biological sample (tissue typing);  
10        and (iii) aid in forensic identification of a biological sample. Some of these applications are described in the subsections, below.

### Chromosome Mapping

15            Once the sequence (or a portion of the sequence) of a gene has been isolated, this sequence can be used to map the location of the gene on a chromosome. This process is called chromosome mapping. Accordingly, portions or fragments of the GPCR<sub>X</sub> sequences, SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or fragments or derivatives thereof, can be used to map the location of the GPCR<sub>X</sub> genes,  
20        respectively, on a chromosome. The mapping of the GPCR<sub>X</sub> sequences to chromosomes is an important first step in correlating these sequences with genes associated with disease.

             Briefly, GPCR<sub>X</sub> genes can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp in length) from the GPCR<sub>X</sub> sequences. Computer analysis of the GPCR<sub>X</sub> sequences can be used to rapidly select primers that do not span more than one exon  
25        in the genomic DNA, thus complicating the amplification process. These primers can then be used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the GPCR<sub>X</sub> sequences will yield an amplified fragment.

             Somatic cell hybrids are prepared by fusing somatic cells from different mammals  
30        (e.g., human and mouse cells). As hybrids of human and mouse cells grow and divide, they gradually lose human chromosomes in random order, but retain the mouse chromosomes. By using media in which mouse cells cannot grow, because they lack a particular enzyme, but in which human cells can, the one human chromosome that contains the gene encoding the needed enzyme will be retained. By using various media, panels of hybrid cell lines can be  
35        established. Each cell line in a panel contains either a single human chromosome or a small

number of human chromosomes, and a full set of mouse chromosomes, allowing easy mapping of individual genes to specific human chromosomes. *See, e.g., D'Eustachio, et al., 1983 Science 220:919-924.* Somatic cell hybrids containing only fragments of human chromosomes can also be produced by using human chromosomes with translocations and deletions.

PCR mapping of somatic cell hybrids is a rapid procedure for assigning a particular sequence to a particular chromosome. Three or more sequences can be assigned per day using a single thermal cycler. Using the GPCRX sequences to design oligonucleotide primers, sub-localization can be achieved with panels of fragments from specific chromosomes.

Fluorescence *in situ* hybridization (FISH) of a DNA sequence to a metaphase chromosomal spread can further be used to provide a precise chromosomal location in one step. Chromosome spreads can be made using cells whose division has been blocked in metaphase by a chemical like colcemid that disrupts the mitotic spindle. The chromosomes can be treated briefly with trypsin, and then stained with Giemsa. A pattern of light and dark bands develops on each chromosome, so that the chromosomes can be identified individually. The FISH technique can be used with a DNA sequence as short as 500 or 600 bases. However, clones larger than 1,000 bases have a higher likelihood of binding to a unique chromosomal location with sufficient signal intensity for simple detection. Preferably 1,000 bases, and more preferably 2,000 bases, will suffice to get good results at a reasonable amount of time. For a review of this technique, *see, Verma, et al., HUMAN CHROMOSOMES: A MANUAL OF BASIC TECHNIQUES* (Pergamon Press, New York 1988).

Reagents for chromosome mapping can be used individually to mark a single chromosome or a single site on that chromosome, or panels of reagents can be used for marking multiple sites and/or multiple chromosomes. Reagents corresponding to noncoding regions of the genes actually are preferred for mapping purposes. Coding sequences are more likely to be conserved within gene families, thus increasing the chance of cross hybridizations during chromosomal mapping.

Once a sequence has been mapped to a precise chromosomal location, the physical position of the sequence on the chromosome can be correlated with genetic map data. Such data are found, *e.g., in McKusick, MENDELIAN INHERITANCE IN MAN*, available on-line through Johns Hopkins University Welch Medical Library). The relationship between genes and disease, mapped to the same chromosomal region, can then be identified through linkage

analysis (co-inheritance of physically adjacent genes), described in, *e.g.*, Egeland, *et al.*, 1987 *Nature*, 325:783-787.

Moreover, differences in the DNA sequences between individuals affected and unaffected with a disease associated with the GPCR<sub>X</sub> gene, can be determined. If a mutation  
5 is observed in some or all of the affected individuals but not in any unaffected individuals, then the mutation is likely to be the causative agent of the particular disease. Comparison of affected and unaffected individuals generally involves first looking for structural alterations in the chromosomes, such as deletions or translocations that are visible from chromosome spreads or detectable using PCR based on that DNA sequence. Ultimately, complete  
10 sequencing of genes from several individuals can be performed to confirm the presence of a mutation and to distinguish mutations from polymorphisms.

### Tissue Typing

The GPCR<sub>X</sub> sequences of the invention can also be used to identify individuals from  
15 minute biological samples. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identification. The sequences of the invention are useful as additional DNA markers for RFLP ("restriction fragment length polymorphisms," described in U.S. Patent No. 5,272,057).

20 Furthermore, the sequences of the invention can be used to provide an alternative technique that determines the actual base-by-base DNA sequence of selected portions of an individual's genome. Thus, the GPCR<sub>X</sub> sequences described herein can be used to prepare two PCR primers from the 5'- and 3'-termini of the sequences. These primers can then be used to amplify an individual's DNA and subsequently sequence it.

25 Panels of corresponding DNA sequences from individuals, prepared in this manner, can provide unique individual identifications, as each individual will have a unique set of such DNA sequences due to allelic differences. The sequences of the invention can be used to obtain such identification sequences from individuals and from tissue. The GPCR<sub>X</sub> sequences of the invention uniquely represent portions of the human genome. Allelic  
30 variation occurs to some degree in the coding regions of these sequences, and to a greater degree in the noncoding regions. It is estimated that allelic variation between individual humans occurs with a frequency of about once per each 500 bases. Much of the allelic variation is due to single nucleotide polymorphisms (SNPs), which include restriction fragment length polymorphisms (RFLPs).

Each of the sequences described herein can, to some degree, be used as a standard against which DNA from an individual can be compared for identification purposes. Because greater numbers of polymorphisms occur in the noncoding regions, fewer sequences are necessary to differentiate individuals. The noncoding sequences can comfortably provide positive individual identification with a panel of perhaps 10 to 1,000 primers that each yield a noncoding amplified sequence of 100 bases. If predicted coding sequences, such as those in SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39 are used, a more appropriate number of primers for positive individual identification would be 500 - 2,000.

### **Predictive Medicine**

The invention also pertains to the field of predictive medicine in which diagnostic assays, prognostic assays, pharmacogenomics, and monitoring clinical trials are used for prognostic (predictive) purposes to thereby treat an individual prophylactically. Accordingly, one aspect of the invention relates to diagnostic assays for determining GPCR<sub>X</sub> protein and/or nucleic acid expression as well as GPCR<sub>X</sub> activity, in the context of a biological sample (*e.g.*, blood, serum, cells, tissue) to thereby determine whether an individual is afflicted with a disease or disorder, or is at risk of developing a disorder, associated with aberrant GPCR<sub>X</sub> expression or activity. The disorders include metabolic disorders, diabetes, obesity, infectious disease, anorexia, cancer-associated cachexia, cancer, neurodegenerative disorders, Alzheimer's Disease, Parkinson's Disorder, immune disorders, and hematopoietic disorders, and the various dyslipidemias, metabolic disturbances associated with obesity, the metabolic syndrome X and wasting disorders associated with chronic diseases and various cancers. The invention also provides for prognostic (or predictive) assays for determining whether an individual is at risk of developing a disorder associated with GPCR<sub>X</sub> protein, nucleic acid expression or activity. For example, mutations in an GPCR<sub>X</sub> gene can be assayed in a biological sample. Such assays can be used for prognostic or predictive purpose to thereby prophylactically treat an individual prior to the onset of a disorder characterized by or associated with GPCR<sub>X</sub> protein, nucleic acid expression, or biological activity.

Another aspect of the invention provides methods for determining GPCR<sub>X</sub> protein, nucleic acid expression or activity in an individual to thereby select appropriate therapeutic or prophylactic agents for that individual (referred to herein as "pharmacogenomics"). Pharmacogenomics allows for the selection of agents (*e.g.*, drugs) for therapeutic or prophylactic treatment of an individual based on the genotype of the individual (*e.g.*, the

genotype of the individual examined to determine the ability of the individual to respond to a particular agent.)

Yet another aspect of the invention pertains to monitoring the influence of agents (e.g., drugs, compounds) on the expression or activity of GPCR<sub>X</sub> in clinical trials.

5           These and other agents are described in further detail in the following sections.

### Diagnostic Assays

10           An exemplary method for detecting the presence or absence of GPCR<sub>X</sub> in a biological sample involves obtaining a biological sample from a test subject and contacting the biological sample with a compound or an agent capable of detecting GPCR<sub>X</sub> protein or nucleic acid (e.g., mRNA, genomic DNA) that encodes GPCR<sub>X</sub> protein such that the presence of GPCR<sub>X</sub> is detected in the biological sample. An agent for detecting GPCR<sub>X</sub> mRNA or genomic DNA is a labeled nucleic acid probe capable of hybridizing to GPCR<sub>X</sub> mRNA or genomic DNA. The nucleic acid probe can be, for example, a full-length GPCR<sub>X</sub> nucleic acid, such as the nucleic acid of SEQ ID NOS:1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37 and 39, or a portion thereof, such as an oligonucleotide of at least 15, 30, 50, 100, 250 or 500 nucleotides in length and sufficient to specifically hybridize under stringent conditions to GPCR<sub>X</sub> mRNA or genomic DNA. Other suitable probes for use in the diagnostic assays of the invention are described herein.

20           An agent for detecting GPCR<sub>X</sub> protein is an antibody capable of binding to GPCR<sub>X</sub> protein, preferably an antibody with a detectable label. Antibodies can be polyclonal, or more preferably, monoclonal. An intact antibody, or a fragment thereof (e.g., Fab or F(ab')<sub>2</sub>) can be used. The term "labeled", with regard to the probe or antibody, is intended to encompass direct labeling of the probe or antibody by coupling (i.e., physically linking) a detectable substance to the probe or antibody, as well as indirect labeling of the probe or antibody by reactivity with another reagent that is directly labeled. Examples of indirect labeling include detection of a primary antibody using a fluorescently-labeled secondary antibody and end-labeling of a DNA probe with biotin such that it can be detected with fluorescently-labeled streptavidin. The term "biological sample" is intended to include tissues, cells and biological fluids isolated from a subject, as well as tissues, cells and fluids present within a subject. That is, the detection method of the invention can be used to detect GPCR<sub>X</sub> mRNA, protein, or genomic DNA in a biological sample *in vitro* as well as *in vivo*. For example, *in vitro* techniques for detection of GPCR<sub>X</sub> mRNA include Northern hybridizations and *in situ* hybridizations. *In vitro* techniques for detection of GPCR<sub>X</sub> protein

include enzyme linked immunosorbent assays (ELISAs), Western blots, immunoprecipitations, and immunofluorescence. *In vitro* techniques for detection of GPCR<sub>X</sub> genomic DNA include Southern hybridizations. Furthermore, *in vivo* techniques for detection of GPCR<sub>X</sub> protein include introducing into a subject a labeled anti-GPCR<sub>X</sub> antibody. For example, the antibody can be labeled with a radioactive marker whose presence and location in a subject can be detected by standard imaging techniques.

In one embodiment, the biological sample contains protein molecules from the test subject. Alternatively, the biological sample can contain mRNA molecules from the test subject or genomic DNA molecules from the test subject. A preferred biological sample is a peripheral blood leukocyte sample isolated by conventional means from a subject.

In another embodiment, the methods further involve obtaining a control biological sample from a control subject, contacting the control sample with a compound or agent capable of detecting GPCR<sub>X</sub> protein, mRNA, or genomic DNA, such that the presence of GPCR<sub>X</sub> protein, mRNA or genomic DNA is detected in the biological sample, and comparing the presence of GPCR<sub>X</sub> protein, mRNA or genomic DNA in the control sample with the presence of GPCR<sub>X</sub> protein, mRNA or genomic DNA in the test sample.

The invention also encompasses kits for detecting the presence of GPCR<sub>X</sub> in a biological sample. For example, the kit can comprise: a labeled compound or agent capable of detecting GPCR<sub>X</sub> protein or mRNA in a biological sample; means for determining the amount of GPCR<sub>X</sub> in the sample; and means for comparing the amount of GPCR<sub>X</sub> in the sample with a standard. The compound or agent can be packaged in a suitable container. The kit can further comprise instructions for using the kit to detect GPCR<sub>X</sub> protein or nucleic acid.

## **Prognostic Assays**

The diagnostic methods described herein can furthermore be utilized to identify subjects having or at risk of developing a disease or disorder associated with aberrant GPCR<sub>X</sub> expression or activity. For example, the assays described herein, such as the preceding diagnostic assays or the following assays, can be utilized to identify a subject having or at risk of developing a disorder associated with GPCR<sub>X</sub> protein, nucleic acid expression or activity. Alternatively, the prognostic assays can be utilized to identify a subject having or at risk for developing a disease or disorder. Thus, the invention provides a method for identifying a disease or disorder associated with aberrant GPCR<sub>X</sub> expression or activity in which a test sample is obtained from a subject and GPCR<sub>X</sub> protein or nucleic acid

(*e.g.*, mRNA, genomic DNA) is detected, wherein the presence of GPCR<sub>X</sub> protein or nucleic acid is diagnostic for a subject having or at risk of developing a disease or disorder associated with aberrant GPCR<sub>X</sub> expression or activity. As used herein, a "test sample" refers to a biological sample obtained from a subject of interest. For example, a test sample can be a biological fluid (*e.g.*, serum), cell sample, or tissue.

Furthermore, the prognostic assays described herein can be used to determine whether a subject can be administered an agent (*e.g.*, an agonist, antagonist, peptidomimetic, protein, peptide, nucleic acid, small molecule, or other drug candidate) to treat a disease or disorder associated with aberrant GPCR<sub>X</sub> expression or activity. For example, such methods can be used to determine whether a subject can be effectively treated with an agent for a disorder. Thus, the invention provides methods for determining whether a subject can be effectively treated with an agent for a disorder associated with aberrant GPCR<sub>X</sub> expression or activity in which a test sample is obtained and GPCR<sub>X</sub> protein or nucleic acid is detected (*e.g.*, wherein the presence of GPCR<sub>X</sub> protein or nucleic acid is diagnostic for a subject that can be administered the agent to treat a disorder associated with aberrant GPCR<sub>X</sub> expression or activity).

The methods of the invention can also be used to detect genetic lesions in an GPCR<sub>X</sub> gene, thereby determining if a subject with the lesioned gene is at risk for a disorder characterized by aberrant cell proliferation and/or differentiation. In various embodiments, the methods include detecting, in a sample of cells from the subject, the presence or absence of a genetic lesion characterized by at least one of an alteration affecting the integrity of a gene encoding an GPCR<sub>X</sub>-protein, or the misexpression of the GPCR<sub>X</sub> gene. For example, such genetic lesions can be detected by ascertaining the existence of at least one of: (i) a deletion of one or more nucleotides from an GPCR<sub>X</sub> gene; (ii) an addition of one or more nucleotides to an GPCR<sub>X</sub> gene; (iii) a substitution of one or more nucleotides of an GPCR<sub>X</sub> gene, (iv) a chromosomal rearrangement of an GPCR<sub>X</sub> gene; (v) an alteration in the level of a messenger RNA transcript of an GPCR<sub>X</sub> gene, (vi) aberrant modification of an GPCR<sub>X</sub> gene, such as of the methylation pattern of the genomic DNA, (vii) the presence of a non-wild-type splicing pattern of a messenger RNA transcript of an GPCR<sub>X</sub> gene, (viii) a non-wild-type level of an GPCR<sub>X</sub> protein, (ix) allelic loss of an GPCR<sub>X</sub> gene, and (x) inappropriate post-translational modification of an GPCR<sub>X</sub> protein. As described herein, there are a large number of assay techniques known in the art which can be used for detecting lesions in an GPCR<sub>X</sub> gene. A preferred biological sample is a peripheral blood leukocyte

sample isolated by conventional means from a subject. However, any biological sample containing nucleated cells may be used, including, for example, buccal mucosal cells.

In certain embodiments, detection of the lesion involves the use of a probe/primer in a polymerase chain reaction (PCR) (*see, e.g.*, U.S. Patent Nos. 4,683,195 and 4,683,202), such as anchor PCR or RACE PCR, or, alternatively, in a ligation chain reaction (LCR) (*see, e.g.*, Landegran, *et al.*, 1988 *Science* 241:1077-1080; and Nakazawa, *et al.*, 1994 *Proc. Natl. Acad. Sci. USA* 91:360-364), the latter of which can be particularly useful for detecting point mutations in the GPCR<sub>X</sub>-gene (*see*, Abravaya, *et al.*, 1995 *Nucl. Acids Res.* 23:675-682). This method can include the steps of collecting a sample of cells from a patient, isolating nucleic acid (*e.g.*, genomic, mRNA or both) from the cells of the sample, contacting the nucleic acid sample with one or more primers that specifically hybridize to an GPCR<sub>X</sub> gene under conditions such that hybridization and amplification of the GPCR<sub>X</sub> gene (if present) occurs, and detecting the presence or absence of an amplification product, or detecting the size of the amplification product and comparing the length to a control sample. It is anticipated that PCR and/or LCR may be desirable to use as a preliminary amplification step in conjunction with any of the techniques used for detecting mutations described herein.

Alternative amplification methods include: self sustained sequence replication (*see*, Guatelli, *et al.*, 1990 *Proc. Natl. Acad. Sci. USA* 87:1874-1878), transcriptional amplification system (*see*, Kwok, *et al.*, 1989 *Proc. Natl. Acad. Sci. USA* 86:1173-1177); Q $\beta$  Replicase (*see*, Lizardi, *et al.*, 1988 *BioTechnology* 6:1197), or any other nucleic acid amplification method, followed by the detection of the amplified molecules using techniques well known to those of skill in the art. These detection schemes are especially useful for the detection of nucleic acid molecules if such molecules are present in very low numbers.

In an alternative embodiment, mutations in an GPCR<sub>X</sub> gene from a sample cell can be identified by alterations in restriction enzyme cleavage patterns. For example, sample and control DNA is isolated, amplified (optionally), digested with one or more restriction endonucleases, and fragment length sizes are determined by gel electrophoresis and compared. Differences in fragment length sizes between sample and control DNA indicates mutations in the sample DNA. Moreover, the use of sequence specific ribozymes (*see, e.g.*, U.S. Patent No. 5,493,531) can be used to score for the presence of specific mutations by development or loss of a ribozyme cleavage site.

In other embodiments, genetic mutations in GPCR<sub>X</sub> can be identified by hybridizing a sample and control nucleic acids, *e.g.*, DNA or RNA, to high-density arrays containing hundreds or thousands of oligonucleotides probes. *See, e.g.*, Cronin, *et al.*, 1996 *Human*



*Mutation* 7:244-255; Kozal, *et al.*, 1996 *Nat. Med.* 2:753-759. For example, genetic mutations in GPCR<sub>X</sub> can be identified in two dimensional arrays containing light-generated DNA probes as described in Cronin, *et al.*, *supra*. Briefly, a first hybridization array of probes can be used to scan through long stretches of DNA in a sample and control to identify base changes between the sequences by making linear arrays of sequential overlapping probes. This step allows the identification of point mutations. This is followed by a second hybridization array that allows the characterization of specific mutations by using smaller, specialized probe arrays complementary to all variants or mutations detected. Each mutation array is composed of parallel probe sets, one complementary to the wild-type gene and the other complementary to the mutant gene.

In yet another embodiment, any of a variety of sequencing reactions known in the art can be used to directly sequence the GPCR<sub>X</sub> gene and detect mutations by comparing the sequence of the sample GPCR<sub>X</sub> with the corresponding wild-type (control) sequence. Examples of sequencing reactions include those based on techniques developed by Maxim and Gilbert, 1977 *Proc. Natl. Acad. Sci. USA* 74:560 or Sanger, 1977 *Proc. Natl. Acad. Sci. USA* 74:5463. It is also contemplated that any of a variety of automated sequencing procedures can be utilized when performing the diagnostic assays (*see, e.g.*, Naeve, *et al.*, 1995 *Biotechniques* 19:448), including sequencing by mass spectrometry (*see, e.g.*, PCT International Publication No. WO 94/16101; Cohen, *et al.*, 1996 *Adv. Chromatography* 36:127-162; and Griffin, *et al.*, 1993 *Appl. Biochem. Biotechnol.* 38:147-159).

Other methods for detecting mutations in the GPCR<sub>X</sub> gene include methods in which protection from cleavage agents is used to detect mismatched bases in RNA/RNA or RNA/DNA heteroduplexes. *See, e.g.*, Myers, *et al.*, 1985 *Science* 230:1242. In general, the art technique of "mismatch cleavage" starts by providing heteroduplexes of formed by hybridizing (labeled) RNA or DNA containing the wild-type GPCR<sub>X</sub> sequence with potentially mutant RNA or DNA obtained from a tissue sample. The double-stranded duplexes are treated with an agent that cleaves single-stranded regions of the duplex such as which will exist due to basepair mismatches between the control and sample strands. For instance, RNA/DNA duplexes can be treated with RNase and DNA/DNA hybrids treated with S<sub>1</sub> nuclease to enzymatically digesting the mismatched regions. In other embodiments, either DNA/DNA or RNA/DNA duplexes can be treated with hydroxylamine or osmium tetroxide and with piperidine in order to digest mismatched regions. After digestion of the mismatched regions, the resulting material is then separated by size on denaturing polyacrylamide gels to determine the site of mutation. *See, e.g.*, Cotton, *et al.*, 1988 *Proc.*

*Natl. Acad. Sci. USA* 85:4397; Saleeba, *et al.*, 1992 *Methods Enzymol.* 217:286-295. In an embodiment, the control DNA or RNA can be labeled for detection.

In still another embodiment, the mismatch cleavage reaction employs one or more proteins that recognize mismatched base pairs in double-stranded DNA (so called "DNA mismatch repair" enzymes) in defined systems for detecting and mapping point mutations in GPCR<sub>X</sub> cDNAs obtained from samples of cells. For example, the mutY enzyme of *E. coli* cleaves A at G/A mismatches and the thymidine DNA glycosylase from HeLa cells cleaves T at G/T mismatches. See, *e.g.*, Hsu, *et al.*, 1994. *Carcinogenesis* 15:1657-1662. According to an exemplary embodiment, a probe based on an GPCR<sub>X</sub> sequence, *e.g.*, a wild-type GPCR<sub>X</sub> sequence, is hybridized to a cDNA or other DNA product from a test cell(s). The duplex is treated with a DNA mismatch repair enzyme, and the cleavage products, if any, can be detected from electrophoresis protocols or the like. See, *e.g.*, U.S. Patent No. 5,459,039.

In other embodiments, alterations in electrophoretic mobility will be used to identify mutations in GPCR<sub>X</sub> genes. For example, single strand conformation polymorphism (SSCP) may be used to detect differences in electrophoretic mobility between mutant and wild type nucleic acids. See, *e.g.*, Orita, *et al.*, 1989 *Proc. Natl. Acad. Sci. USA*: 86:2766; Cotton, 1993 *Mutat. Res.* 285:125-144; Hayashi, 1992 *Genet. Anal. Tech. Appl.* 9:73-79. Single-stranded DNA fragments of sample and control GPCR<sub>X</sub> nucleic acids will be denatured and allowed to renature. The secondary structure of single-stranded nucleic acids varies according to sequence, the resulting alteration in electrophoretic mobility enables the detection of even a single base change. The DNA fragments may be labeled or detected with labeled probes. The sensitivity of the assay may be enhanced by using RNA (rather than DNA), in which the secondary structure is more sensitive to a change in sequence. In one embodiment, the subject method utilizes heteroduplex analysis to separate double stranded heteroduplex molecules on the basis of changes in electrophoretic mobility. See, *e.g.*, Keen, *et al.*, 1991 *trends Genet.* 7:5.

In yet another embodiment, the movement of mutant or wild-type fragments in polyacrylamide gels containing a gradient of denaturant is assayed using denaturing gradient gel electrophoresis (DGGE). See, *e.g.*, Myers, *et al.*, 1985 *Nature* 313:495. When DGGE is used as the method of analysis, DNA will be modified to insure that it does not completely denature, for example by adding a GC clamp of approximately 40 bp of high-melting GC-rich DNA by PCR. In a further embodiment, a temperature gradient is used in place of a denaturing gradient to identify differences in the mobility of control and sample DNA. See, *e.g.*, Rosenbaum and Reissner, 1987 *Biophys. Chem.* 265:12753.

Examples of other techniques for detecting point mutations include, but are not limited to, selective oligonucleotide hybridization, selective amplification, or selective primer extension. For example, oligonucleotide primers may be prepared in which the known mutation is placed centrally and then hybridized to target DNA under conditions that permit hybridization only if a perfect match is found. *See, e.g., Saiki, et al., 1986 Nature 324:163; Saiki, et al., 1989 Proc. Natl. Acad. Sci. USA 86:6230.* Such allele specific oligonucleotides are hybridized to PCR amplified target DNA or a number of different mutations when the oligonucleotides are attached to the hybridizing membrane and hybridized with labeled target DNA.

Alternatively, allele specific amplification technology that depends on selective PCR amplification may be used in conjunction with the instant invention. Oligonucleotides used as primers for specific amplification may carry the mutation of interest in the center of the molecule (so that amplification depends on differential hybridization; *see, e.g., Gibbs, et al., 1989 Nucl. Acids Res. 17:2437-2448*) or at the extreme 3'-terminus of one primer where, under appropriate conditions, mismatch can prevent, or reduce polymerase extension (*see, e.g., Prossner, 1993 Tibtech. 11:238*). In addition it may be desirable to introduce a novel restriction site in the region of the mutation to create cleavage-based detection. *See, e.g., Gasparini, et al., 1992 Mol. Cell Probes 6:1.* It is anticipated that in certain embodiments amplification may also be performed using *Taq* ligase for amplification. *See, e.g., Barany, 1991 Proc. Natl. Acad. Sci. USA 88:189.* In such cases, ligation will occur only if there is a perfect match at the 3'-terminus of the 5' sequence, making it possible to detect the presence of a known mutation at a specific site by looking for the presence or absence of amplification.

The methods described herein may be performed, for example, by utilizing pre-packaged diagnostic kits comprising at least one probe nucleic acid or antibody reagent described herein, which may be conveniently used, *e.g.,* in clinical settings to diagnose patients exhibiting symptoms or family history of a disease or illness involving an GPCR<sub>X</sub> gene.

Furthermore, any cell type or tissue, preferably peripheral blood leukocytes, in which GPCR<sub>X</sub> is expressed may be utilized in the prognostic assays described herein. However, any biological sample containing nucleated cells may be used, including, for example, buccal mucosal cells.

## Pharmacogenomics

Agents, or modulators that have a stimulatory or inhibitory effect on GPCR<sub>X</sub> activity (e.g., GPCR<sub>X</sub> gene expression), as identified by a screening assay described herein can be administered to individuals to treat (prophylactically or therapeutically) disorders (The disorders include metabolic disorders, diabetes, obesity, infectious disease, anorexia, cancer-associated cachexia, cancer, neurodegenerative disorders, Alzheimer's Disease, Parkinson's Disorder, immune disorders, and hematopoietic disorders, and the various dyslipidemias, metabolic disturbances associated with obesity, the metabolic syndrome X and wasting disorders associated with chronic diseases and various cancers.) In conjunction with such treatment, the pharmacogenomics (*i.e.*, the study of the relationship between an individual's genotype and that individual's response to a foreign compound or drug) of the individual may be considered. Differences in metabolism of therapeutics can lead to severe toxicity or therapeutic failure by altering the relation between dose and blood concentration of the pharmacologically active drug. Thus, the pharmacogenomics of the individual permits the selection of effective agents (e.g., drugs) for prophylactic or therapeutic treatments based on a consideration of the individual's genotype. Such pharmacogenomics can further be used to determine appropriate dosages and therapeutic regimens. Accordingly, the activity of GPCR<sub>X</sub> protein, expression of GPCR<sub>X</sub> nucleic acid, or mutation content of GPCR<sub>X</sub> genes in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual.

Pharmacogenomics deals with clinically significant hereditary variations in the response to drugs due to altered drug disposition and abnormal action in affected persons. See *e.g.*, Eichelbaum, 1996 *Clin. Exp. Pharmacol. Physiol.*, 23:983-985; Linder, 1997 *Clin. Chem.*, 43:254-266. In general, two types of pharmacogenetic conditions can be differentiated. Genetic conditions transmitted as a single factor altering the way drugs act on the body (altered drug action) or genetic conditions transmitted as single factors altering the way the body acts on drugs (altered drug metabolism). These pharmacogenetic conditions can occur either as rare defects or as polymorphisms. For example, glucose-6-phosphate dehydrogenase (G6PD) deficiency is a common inherited enzymopathy in which the main clinical complication is hemolysis after ingestion of oxidant drugs (anti-malarials, sulfonamides, analgesics, nitrofurans) and consumption of fava beans.

As an illustrative embodiment, the activity of drug metabolizing enzymes is a major determinant of both the intensity and duration of drug action. The discovery of genetic

polymorphisms of drug metabolizing enzymes (*e.g.*, N-acetyltransferase 2 (NAT 2) and cytochrome P450 enzymes CYP2D6 and CYP2C19) has provided an explanation as to why some patients do not obtain the expected drug effects or show exaggerated drug response and serious toxicity after taking the standard and safe dose of a drug. These polymorphisms are expressed in two phenotypes in the population, the extensive metabolizer (EM) and poor metabolizer (PM). The prevalence of PM is different among different populations. For example, the gene coding for CYP2D6 is highly polymorphic and several mutations have been identified in PM, which all lead to the absence of functional CYP2D6. Poor metabolizers of CYP2D6 and CYP2C19 quite frequently experience exaggerated drug response and side effects when they receive standard doses. If a metabolite is the active therapeutic moiety, PM show no therapeutic response, as demonstrated for the analgesic effect of codeine mediated by its CYP2D6-formed metabolite morphine. At the other extreme are the so called ultra-rapid metabolizers who do not respond to standard doses. Recently, the molecular basis of ultra-rapid metabolism has been identified to be due to CYP2D6 gene amplification.

Thus, the activity of GPCR<sub>X</sub> protein, expression of GPCR<sub>X</sub> nucleic acid, or mutation content of GPCR<sub>X</sub> genes in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual. In addition, pharmacogenetic studies can be used to apply genotyping of polymorphic alleles encoding drug-metabolizing enzymes to the identification of an individual's drug responsiveness phenotype. This knowledge, when applied to dosing or drug selection, can avoid adverse reactions or therapeutic failure and thus enhance therapeutic or prophylactic efficiency when treating a subject with an GPCR<sub>X</sub> modulator, such as a modulator identified by one of the exemplary screening assays described herein.

### Monitoring of Effects During Clinical Trials

Monitoring the influence of agents (*e.g.*, drugs, compounds) on the expression or activity of GPCR<sub>X</sub> (*e.g.*, the ability to modulate aberrant cell proliferation and/or differentiation) can be applied not only in basic drug screening, but also in clinical trials. For example, the effectiveness of an agent determined by a screening assay as described herein to increase GPCR<sub>X</sub> gene expression, protein levels, or upregulate GPCR<sub>X</sub> activity, can be monitored in clinical trials of subjects exhibiting decreased GPCR<sub>X</sub> gene expression, protein levels, or downregulated GPCR<sub>X</sub> activity. Alternatively, the effectiveness of an agent determined by a screening assay to decrease GPCR<sub>X</sub> gene expression, protein levels, or

downregulate GPCR<sub>X</sub> activity, can be monitored in clinical trials of subjects exhibiting increased GPCR<sub>X</sub> gene expression, protein levels, or upregulated GPCR<sub>X</sub> activity. In such clinical trials, the expression or activity of GPCR<sub>X</sub> and, preferably, other genes that have been implicated in, for example, a cellular proliferation or immune disorder can be used as a  
5 "read out" or markers of the immune responsiveness of a particular cell.

By way of example, and not of limitation, genes, including GPCR<sub>X</sub>, that are modulated in cells by treatment with an agent (*e.g.*, compound, drug or small molecule) that modulates GPCR<sub>X</sub> activity (*e.g.*, identified in a screening assay as described herein) can be identified. Thus, to study the effect of agents on cellular proliferation disorders, for example,  
10 in a clinical trial, cells can be isolated and RNA prepared and analyzed for the levels of expression of GPCR<sub>X</sub> and other genes implicated in the disorder. The levels of gene expression (*i.e.*, a gene expression pattern) can be quantified by Northern blot analysis or RT-PCR, as described herein, or alternatively by measuring the amount of protein produced, by one of the methods as described herein, or by measuring the levels of activity of GPCR<sub>X</sub>  
15 or other genes. In this manner, the gene expression pattern can serve as a marker, indicative of the physiological response of the cells to the agent. Accordingly, this response state may be determined before, and at various points during, treatment of the individual with the agent.

In one embodiment, the invention provides a method for monitoring the effectiveness of treatment of a subject with an agent (*e.g.*, an agonist, antagonist, protein, peptide,  
20 peptidomimetic, nucleic acid, small molecule, or other drug candidate identified by the screening assays described herein) comprising the steps of (i) obtaining a pre-administration sample from a subject prior to administration of the agent; (ii) detecting the level of expression of an GPCR<sub>X</sub> protein, mRNA, or genomic DNA in the preadministration sample; (iii) obtaining one or more post-administration samples from the subject; (iv) detecting the  
25 level of expression or activity of the GPCR<sub>X</sub> protein, mRNA, or genomic DNA in the post-administration samples; (v) comparing the level of expression or activity of the GPCR<sub>X</sub> protein, mRNA, or genomic DNA in the pre-administration sample with the GPCR<sub>X</sub> protein, mRNA, or genomic DNA in the post administration sample or samples; and (vi) altering the administration of the agent to the subject accordingly. For example, increased administration  
30 of the agent may be desirable to increase the expression or activity of GPCR<sub>X</sub> to higher levels than detected, *i.e.*, to increase the effectiveness of the agent. Alternatively, decreased administration of the agent may be desirable to decrease expression or activity of GPCR<sub>X</sub> to lower levels than detected, *i.e.*, to decrease the effectiveness of the agent.

## Methods of Treatment

The invention provides for both prophylactic and therapeutic methods of treating a subject at risk of (or susceptible to) a disorder or having a disorder associated with aberrant GPCR<sub>X</sub> expression or activity. The disorders include cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, atrial septal defect (ASD), atrioventricular (A-V) canal defect, ductus arteriosus, pulmonary stenosis, subaortic stenosis, ventricular septal defect (VSD), valve diseases, tuberous sclerosis, scleroderma, obesity, transplantation, adrenoleukodystrophy, congenital adrenal hyperplasia, prostate cancer, neoplasm; adenocarcinoma, lymphoma, uterus cancer, fertility, hemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, immunodeficiencies, graft versus host disease, AIDS, bronchial asthma, Crohn's disease; multiple sclerosis, treatment of Albright hereditary osteodystrophy, and other diseases, disorders and conditions of the like. These methods of treatment will be discussed more fully, below.

## Disease and Disorders

Diseases and disorders that are characterized by increased (relative to a subject not suffering from the disease or disorder) levels or biological activity may be treated with Therapeutics that antagonize (*i.e.*, reduce or inhibit) activity. Therapeutics that antagonize activity may be administered in a therapeutic or prophylactic manner. Therapeutics that may be utilized include, but are not limited to: (i) an aforementioned peptide, or analogs, derivatives, fragments or homologs thereof; (ii) antibodies to an aforementioned peptide; (iii) nucleic acids encoding an aforementioned peptide; (iv) administration of antisense nucleic acid and nucleic acids that are "dysfunctional" (*i.e.*, due to a heterologous insertion within the coding sequences of coding sequences to an aforementioned peptide) that are utilized to "knockout" endogenous function of an aforementioned peptide by homologous recombination (*see, e.g.*, Capecchi, 1989. *Science* 244: 1288-1292); or (v) modulators (*i.e.*, inhibitors, agonists and antagonists, including additional peptide mimetic of the invention or antibodies specific to a peptide of the invention) that alter the interaction between an aforementioned peptide and its binding partner. Diseases and disorders that are characterized by decreased (relative to a subject not suffering from the disease or disorder) levels or biological activity may be treated with therapeutics that increase (*i.e.*, are agonists to) activity. Therapeutics that upregulate activity may be administered in a therapeutic or prophylactic manner. Therapeutics that may be utilized include, but are not limited to, an aforementioned peptide, or analogs, derivatives, fragments or homologs thereof; or an agonist

that increases bioavailability. Increased or decreased levels can be readily detected by quantifying peptide and/or RNA, by obtaining a patient tissue sample (*e.g.*, from biopsy tissue) and assaying it *in vitro* for RNA or peptide levels, structure and/or activity of the expressed peptides (or mRNAs of an aforementioned peptide). Methods that are well-known within the art include, but are not limited to, immunoassays (*e.g.*, by Western blot analysis, immunoprecipitation followed by sodium dodecyl sulfate (SDS) polyacrylamide gel electrophoresis, immunocytochemistry, etc.) and/or hybridization assays to detect expression of mRNAs (*e.g.*, Northern assays, dot blots, *in situ* hybridization, and the like).

## 10 Prophylactic Methods

In one aspect, the invention provides a method for preventing, in a subject, a disease or condition associated with an aberrant GPCR<sub>X</sub> expression or activity, by administering to the subject an agent that modulates GPCR<sub>X</sub> expression or at least one GPCR<sub>X</sub> activity. Subjects at risk for a disease that is caused or contributed to by aberrant GPCR<sub>X</sub> expression or activity can be identified by, for example, any or a combination of diagnostic or prognostic assays as described herein. Administration of a prophylactic agent can occur prior to the manifestation of symptoms characteristic of the GPCR<sub>X</sub> aberrancy, such that a disease or disorder is prevented or, alternatively, delayed in its progression. Depending upon the type of GPCR<sub>X</sub> aberrancy, for example, an GPCR<sub>X</sub> agonist or GPCR<sub>X</sub> antagonist agent can be used for treating the subject. The appropriate agent can be determined based on screening assays described herein. The prophylactic methods of the invention are further discussed in the following subsections.

## Therapeutic Methods

Another aspect of the invention pertains to methods of modulating GPCR<sub>X</sub> expression or activity for therapeutic purposes. The modulatory method of the invention involves contacting a cell with an agent that modulates one or more of the activities of GPCR<sub>X</sub> protein activity associated with the cell. An agent that modulates GPCR<sub>X</sub> protein activity can be an agent as described herein, such as a nucleic acid or a protein, a naturally-occurring cognate ligand of an GPCR<sub>X</sub> protein, a peptide, an GPCR<sub>X</sub> peptidomimetic, or other small molecule. In one embodiment, the agent stimulates one or more GPCR<sub>X</sub> protein activity. Examples of such stimulatory agents include active GPCR<sub>X</sub> protein and a nucleic acid molecule encoding GPCR<sub>X</sub> that has been introduced into the cell. In another embodiment, the agent inhibits one or more GPCR<sub>X</sub> protein activity. Examples of



such inhibitory agents include antisense GPCR<sub>X</sub> nucleic acid molecules and anti-GPCR<sub>X</sub> antibodies. These modulatory methods can be performed *in vitro* (e.g., by culturing the cell with the agent) or, alternatively, *in vivo* (e.g., by administering the agent to a subject). As such, the invention provides methods of treating an individual afflicted with a disease or disorder characterized by aberrant expression or activity of an GPCR<sub>X</sub> protein or nucleic acid molecule. In one embodiment, the method involves administering an agent (e.g., an agent identified by a screening assay described herein), or combination of agents that modulates (e.g., up-regulates or down-regulates) GPCR<sub>X</sub> expression or activity. In another embodiment, the method involves administering an GPCR<sub>X</sub> protein or nucleic acid molecule as therapy to compensate for reduced or aberrant GPCR<sub>X</sub> expression or activity.

Stimulation of GPCR<sub>X</sub> activity is desirable in situations in which GPCR<sub>X</sub> is abnormally downregulated and/or in which increased GPCR<sub>X</sub> activity is likely to have a beneficial effect. One example of such a situation is where a subject has a disorder characterized by aberrant cell proliferation and/or differentiation (e.g., cancer or immune associated disorders). Another example of such a situation is where the subject has a gestational disease (e.g., preclampsia).

#### **Determination of the Biological Effect of the Therapeutic**

In various embodiments of the invention, suitable *in vitro* or *in vivo* assays are performed to determine the effect of a specific Therapeutic and whether its administration is indicated for treatment of the affected tissue.

In various specific embodiments, *in vitro* assays may be performed with representative cells of the type(s) involved in the patient's disorder, to determine if a given Therapeutic exerts the desired effect upon the cell type(s). Compounds for use in therapy may be tested in suitable animal model systems including, but not limited to rats, mice, chicken, cows, monkeys, rabbits, and the like, prior to testing in human subjects. Similarly, for *in vivo* testing, any of the animal model system known in the art may be used prior to administration to human subjects.

#### **Prophylactic and Therapeutic Uses of the Compositions of the Invention**

The GPCR<sub>X</sub> nucleic acids and proteins of the invention have prophylactic and therapeutic applications implicated in a variety of disorders, not limited to: metabolic disorders, diabetes, obesity, infectious diseases,

associated cancer, neurodegenerative disorders, Alzheimer's disease, Parkinson's disorder, immune disorders, hematopoietic disorders, and the various dyslipidemias, metabolic disturbances associated with obesity, the metabolic syndrome X and wasting disorders associated with chronic diseases and various cancers.

5           As an example, a cDNA encoding the GPCR<sub>X</sub> protein of the invention may be useful in gene therapy, and the protein may be useful when administered to a subject in need thereof. By way of non-limiting example, the compositions of the invention will have efficacy for treatment of patients suffering from: metabolic disorders, diabetes, obesity, infectious disease, anorexia, cancer-associated cachexia, cancer, neurodegenerative disorders, Alzheimer's disease, Parkinson's disorder, immune disorders, hematopoietic disorders, and  
10           the various dyslipidemias.

          Both the novel nucleic acid encoding the GPCR<sub>X</sub> protein, and the GPCR<sub>X</sub> protein of the invention, or fragments thereof, may also be useful in diagnostic applications, wherein the presence or amount of the nucleic acid or the protein are to be assessed. A further use could  
15           be as an anti-bacterial molecule (*i.e.*, some peptides have been found to possess anti-bacterial properties). These materials are further useful in the generation of antibodies which immunospecifically-bind to the novel substances of the invention for use in therapeutic or diagnostic methods.

          The invention will be further described in the following examples, which do not limit  
20           the scope of the invention described in the claims.



Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAP00112_C	3/4	GAAATTATGAGAAGAAACCGTACATTGGTGACTGAGTTCAATCTCCTGGGACTGG CCAATCACTGGGAATTACAGAAATTTCCCTCTTCACGCTGTTTCTCACCATTTACATG GTCACGGTGGCAGGAAATCTTGGCATGATTGCCCTCATCCAGGCCAACCCCGGC TCCACACGCCCATGTACTTTTCTGTAGCAACTTATCCCTTTGTGGAATCTGTGCTTC TCTTCCAAATGTGACTCCCAAGGATGCTGGAGATTTTCCCTTTCAGAGAGAAAGCA TTTCCCTATCCTGCCGCTTGTGCAGTGTACCTTTTATCACCTTGGTCCACGTT GAGCTCTACATCCTGGCTGTGATGGCCCTTTGACCGGTACATGGCCATCTGCAACC CTCTGCTTTATGGCAGCAGAAATGTCCAAGAGCGTGTGCTCTTTCCTCATCACAGT GCTTTATGTGATGGAGCACTCACTGGCCTGATGGAGACTATGTGGACCTACAAC CTAGCCTTCTGTGGCCCCAGTGAAATTAATCACTTCTACTGTGTGGACCCACCACT GATTAAGCTGGCTTGTCTGACACCTACAACAGGAGGTGTCAATGTTTGTGTG GCTGGTTTCAACTTCACCTTATCCTCTCCTTATCATCCTCATTTCCCTATCTCTACATA TTTCTGCCACCCCTAAGGATCTGCTCTACAGAAAGGCAGGCACAAAGCTTTTCTA CCTGTGGCTCCCATCTGACAGCCGTTACTATTTCTATTTCAGCTCTTTTCTTCATG TATCTCAGACGTCCATCAGAAAGAGTCCATGGAGCAGGGGAAATGGTAGCTGTA TTTTATACCACTGTAAATCCCCATGTTGAATCCCATGATCTACAGTCTGAGGAAACA AAGATGTGAAAGAGGCATTATGCAAGAAAGAACTGTTCAAAAGAAATGTTTCTTAA ATAAACATTACTACTGATTTTTT	MRRNRTLVTFFILLGLAN HWELQIFLFTFLTYYMV TVAGNLGMIALIQANPRL HTPMYFFLSNLSFVDLCF SSNVTPRMLEIFLSEKSI SYPARLVQCYLFITLVHV ELYILAVMAFDRYMAICN PLLYGSRMSKSVCSFLIT VLYVYGALTGLMETMW TYNLAFCGPSEINHFCV DPPLIKLACSDTYNKEVS MFVVAGFNFTYPLLIILIS YLYIFPATLRICSTEGRHK AFSTCGSHLTAVTIFYSL FFMYLRRPSEESMEQ GK MVAVFYTTVIPMLNPMI YSLRNKDVKEALCKELF KRKLFSK

Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAC025115_A_5/6		CATGGAAGGGGAAATTGGACATTGGTGACTGAGTTTATTCTTGTGGGGATACCA ACCACAGAGCCCTTGGGGCCCTCCTCTTTTATCAGCCTATTGTGTGACAGTCCCT TGGAAACACCCCTTATTATATCCTGATTCTTGTGATTACAGGCTCCACTCACCCA TGTATTTCTTCTCAGCAATCTCTCTTTTCAGTGAAACAATTAAACCATAAACCTGTGCT GTTCCCTAAGATGCTGGAGGGCTTCCCGTCGGAAAGGAAGAGCATCACAAAGTGGC GAAATGCTCTGCACAGTCCCTATTCTCTATTTCTTCCGGATGCACGTGATTATCC TTTTGTCTGTCATGTCCCTATGACCGCTATGTGGCCTATTGCGATCCTCTTCAGTACC CTGCAATTATGACCAGCTCACTCTGTGCCACCTCGTCACTCTCTCCTGGGTGGGT GGCTTTCTCCTCATGTCCCATCCACCATCCTCAAGGCAGGACTGCCACACTGTG GTCCCAACGTGATTGAGCACTTTTCTGTGACAGCGCCCTCTCCTCCACCTGGCC TGTGCTGACATTCTGCTATTGAGCTGTGGACTTTCTCAGCTCACTGGTCTCTGAT CCTCAGCTCCCTCTCACTCACAGTGGTCTCTATGTTTACATCATCTCCACCATTC TGAAGATACCTCAGGCCAAGGTCAACGCAAGCCTTGGCACCTGTGCCCTCTCA CTTACGGTGTCTCCGTGGGCTATGGGATCTCCATCTTGTCTATGTTCAACCCCT CACAGAAGAGCAGCCTGCACCTCAACAAGATCCTCTTTATCCTCTCCAGCATCAT CACACCCCTCCTGAATCCCTTCGTCTTCAGTCTGTGGAATGAACCCATGAAAGAT GCACTGAAGGACCGCCTCGGCCGGAGGACAGAGCTTGCTCAAAAGGGGTAAAGTCC ACATGAGGATTCTCTGAGAAT	MERGNWTLVTEFLVGP TTRALGGLFLSAYLVTV LGNTLIILILVDYRLHSP MYFFLSNLSFSETLTITCA VPKMLEGFPSEKKSITSG ECSAQSYFYFLSGCTEFIP FAVMSYDRYVVAICSPLOY PAIMTSSSLCAHLVILSWV GGFLMLPSTILKAGLPH CGPNVIEHFFCDSAPLLH LACADIRAIELLDLFLSLV LILSSLSLTVVSVYIIST LKIPSGQGQRKAFATCAS HFTVSVVGYGISIFVYVH PSQKSSLHLNKLILFILSSIIT PLLNPFVFSLWNEPMKD ALKDASAGGQSLKGV ST

Table 1

Acc. No.	SEQ ID NO (Nuel) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAP002512_C_7/8		ACCAATGCGTCTCATAAGAGATGAAGAAATGTCCAGAAGAACTATACTGAAC GACAGAAATTTGTTCTCTGGGTCTAACAAAGCCGTCCAGAGCTGCGAGTTGCTTTC TTGGCACTGTTCCCTTTTGTCTACATAGCCACTGTGTAGGAAACTTGGGGATGA TTATTTTAATCAAAAGTTGATTCTCGACTTCACACTCCCATGGAAATTTTCTCTCC AGTTGTCCATTCTAGATCTGTGTTTCTCCACAAAATTCACCTCCCAAAATGCTAGA AAATTTCTTATCAGAGAAGAAAGACCAATTCCTATGCAGGTTGTTTGATGCAGTGC TATGTTGTCAATTGCTGTGGTCCTTGACAGACACTGCATGTTGGCAGTCATGGCAT ATGACCGCTATATGGCCATCTGTAATCCATTGCTCTACAGTAGCAAAATGTCCCA AGGTGTTGTGTCCACCTGGTCATTGTCCCTTATGTCTATGGCTTCTCTCAGTG TGATGAAACCTTAAGGACCTACAACCTCTCCTTCTGTGAACAAATGAAATCAA CCATTTCTACTGTGCTGATCCTCCTCTTATCAAACTGGCATGCTCTGACACGTACA GCAAGGAGCTGTCCATGTACATAGTAGCCGGCTACAGCAACGTCCTCAGTCTCTCT GATCATCTCACATCCTACATGTTTCATCCTGTGCTATCCTCAGAAAGCCATTCTG CAGAGGGAAGGAAAGCTTTTCCACATGTGTTCCACCTGACAGTTGTGTCAC AATCTTCTATGGAACCTCTTCTGCATGCATTTGAGACGTCCCAACAGACGATCC GTGGAGCAGGGGAAATGGTGGCTGTGTTTACACCAAGTGTATCTCATGTCTGA ACTCCATGATCTATGGCCTCAGGAACAAAGGATGTGAAAGAGCGGTTGAAAAAAG CAATAGGAAACAAACATTGGGAAATAAAATGCTAAGCTATCATTA	MRLRDEEMSRRNRYTELT EFVLLGLTSRPELRVAFI ALFLFVYIATVVGNLGMI ILIKVDSRLHTPMEFFLSS LSILDLCFSTNFTPKMLEN FLSEKKTISYAGCLMQCY VVIAVVLAEHCMLAVMA YDRYMAICNPLLYSSKM SQGVCVHLVIVPYVYGFL LSVMETLR.TYNLSFCGTN EINHFYCADPPLIKLACSD TYSKELSMYIVAGYSNV QSLILTSYMFILVAILRS HSAEGRKKAFSTCGSHLT VVTIFYGTLFCMHLLRRPT DESVEQGMVAVFYTTV ILMLNSMIYGLRNKDVKE ALKKAIGKQTLGK

Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAP002358_B	9/10	TGCAATGACTGGGAAAGGAAACAGTACGAGAATTACAAAAGTTCAATCTCTTGGG ATTCTCTGAATTTCCAAAGAACCCCTATTTCTCTTTTCAATAATTCCTAGGGATCT ACCTCCTGACAGTGTCTGGAACATAAACCTCATCACCCCTATCAGGACGGACTC CCATCTGCATACACCTATGTACTTTTCTTCCCTTAGTAATCTGCTGTTCTGGACATCT GCTATGTTTCCACTATAGCCCCCAAGATGCTCTCAGACTTCTTCAAGAAGCATAA ATTCACTCTCTTATGGGGTGCAGTATGCAGTACTTTTCTCTCTAGCCTAGGTC TAACTGAGTGTCTTCTGGCAGCCATGGCTTATGATCGATATGCTGCCATTG CAACCCCTCTGCTACAGGGCCATCATGTTTCCCACCCCTCTGCGTGCAGATGGTG GCAGGATCTTGATAACTGGATTCTTAGGCTCATTTATCCAACTCTGTGCTTGTCT TCAGTCCATTCTGTGGGCCAAATGTCAACAACCATTTCTTCTGTGATCTGCCCC AGCTGCTGATCTATCCTGTCTGACACCTTTTCTTCAAGTCATGACCTCTGTT CTCACAGTGATCTTGGACTCACGTCGTCTTAGTTATCATGATATCTTATGGTTA TATCATTGCCACCATCTGAAGATCACCTCAGCTGAAGCAGAGCCAAATCTTTC AACACTTGCTTCTCACCTTACAGCAGTGATCCCTTTTCTTGGCTCAGGTATCTT TGTTTATATGATCCTAATGCTGGTGATTCCTTGAGCCAAACAAAGTTGGCATCA GTCTTATACACAGTTACAATCCCCCATGTTAAATCCAGTGATCTACAGCCTGAGGA ACAAAGGAAATCAAAGATGCTCTAAACAGATGGAAGAAAGAGAAATCTTCTCCTGGT GTTATGGAATGAAATAATGGAAATTTATTTTCAGAT	MTGERNSTRIITKFIILGFS EFPKNPIFLSIFLGIYLLT VSWNINLITLIRTDShLHT PMYFFLSNLSFLDICYVST IAPKMLSDFFKKHKFISF MGCSMQYFFFSLSLGLTEC CLLAAMAYDRYAAICNP LLYRAIMFPTLCVQMVA GSCITGFLGSFIQLCALLQ LHFCGPNVINHFFCDLPQ LLJLSCSDTFFQVMTSVL TVIFGLTSVLVIMISYGYII ATILKITSAEGRAKSFNTC ASHLTAVILFFGSGIFVY MYPNAGDSLSQNKLASV LYTVTIPMLNPVIYSLRN KEIKDALNRWKKRIFSW CYGMK

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Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAP002418_C	11/12	GCTTTCACCTCCGTGGACATGGAAGTGGGAAATTGCACCATCCTGACTGAATTCA TCTTGTGGGTTTCTCAGCAGATTCCAGTGGCAGCCGATTCTATTGGAGTGT CTGATGCTCTATTGATAACCTTGTCAAGAAACATGACCTGGTTATCTTAATCCG AACTGATCCCACTTGCATACACCTATGTACTTTTTCATTGGCAATCTGTCTTTT TGGATTTCTGGTATACCTCTGTGTATACCCCAAAATCCTGGCCAGTTGTCTCA GAAGATAAGCGCATTTCTTGGCTGGATGTGGGCTCAGCTGTTTTTTCCTGTG TTGTAGCCTACACTGAATGCTATCTCCTGGCAGCCATGGCATATGACCGCCATGC AGCAATTTGTAAACCAATTGCTTTATTCAGGTACCATGTCCACCCCTCTGTACTG GGCTTGTGCTGGCTCCTACATAGGAGGATTTTGTGAATGCCATAGCCCATACTGC CAATACATTCGGCTGCATTTTGTGGTAAATATCATTGACCACTTTTCTGTG ATGCACCAACATTGGTAAATGTCTGTACAGACACCCAGGTCTACGAAAAAGT CCTGCTTGGTGGTGGCTTACAGCACTCTCCAGCATTTCTGTATCCTGATTT CCTATGTCAACATCCTCCTGGCTATCCTGAGAAATCCACTCAGCTTCAGGAAGACA CAAGGCATTCTCCACCTGTGCTTCCACCTCAGTCCACCTACTCCCTAGAGGGACAA CATTGTTGTTATGTATTCAAGGCCTAGTCCACCTACTCCCTAGAGGGACAA AGTAGCTGCTGTTCTACACCGTGATCAACCCCACTGCTCAACCCCTCATCTATA GCCTGAGAAACAAAGATATCAAAAGAGGCCCTTCAGGAAAGCAACACAGACTATAC AACCACAAACATGAAG	MEVGNCTILTEFILLGFSA DSQWQPILFGVFLMLYLI TLSGNMTLVILIRTDShL HTPMYFFIGNLSFLDFWY TSVYTPKILASCVSEDKRI SLAGCGAQLFFSCVVA YT ECYLLAAMAYDRHAAIC NPLLYSGTMSTALCTGLV AGSYIGGFLNAIAHTANT FRLHFCGKNIIDHFFCDAP PLVKMSCTDTRVYEKVL LGVVGFTALSSILAILSY VNILLAILRHSAASGRHKA FSTCASHLISVMLFYGSLL FMYSRPSSTYSLERDKVA ALFYTVINPLLNPLIYSLR NKDIKEAFRKATQTIQPO T



Table 1

Acc. No.	SEQ ID NO (NucI) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAP002512_B	13/14	TAATAATCATGTTGCTATTTGTAATTGTAATCCAATATATATGAAAAGTTCCTTTCTT CCACCGAAGGAAATTATGAGAAAGAACTGCACGTTGGTGACTGAGTTCATTCTCC TGGGACTGACCAAGTCGCCGGGAATTACAAATTCTCCTCTTCACGCTGTTCTGCGC CATTACATGGTCACGGTGGCAGGGAACCTTGGCATGATTGTCTCATCCAGGCC AACGCCTGGCTCCACATGCCCATGTACTTTTCTGAGCCACTTATCCTTCGTGGA TCTGTGCTTCTCTCCAAATGTGACTCCAAAGATGCTGGAGATTTTCTTTCAGAGA AGAAAAGCATTTCTATCCTGCTGCTGTGTGTCAGTGTACCTTTTATCGCCTTG GTCCATGTTGAGATCTACATCCTGGCTGTGATGGCCTTGGACCGGTACATGGCCA TCTGCAACCCCTCTGCTTATGGCAGCAGAAATGTCCAAGAGTGTGTCTCCTTCCTC ATCACGGTGCCTTATGTGATGGAGCGCTCACTGGCCTGATGGAGACCATGTGGA CCTACAACCTAGCCTTCTGTGGCCCCAAATGAAATTAATCACTTCTACTGTGCGGA CCCACCACTGATTAAAGCTGGCTGTCTGACACCTACAACAAGGAGTTGTCAATG TTTATTGTGGCTGGTGAACCTTTCTTTTCTCTCTTCATCATATGTATTCTCTAC CTTTACATTTTCCCTGCTATTTAAAGATTCTGCTCTACAGAGGCGGCAAAAAG CTTTTCTACCTGTGGCTCCCATCTGACAGCTGTCACTATATTCTATGCCACCCCT TTCTTCATGTATCTCAGACCCCTCAAGGAATCTGTTGAACAGGGTAAATGG TAGCTGATTTTATACCACAGTAATCCCTATGCTGAACCTTATAATTTATAGCCTT AGAAATAAAAATGTAAAGAAGCATTAATCAAAAGAGCTGTCAATGAAGATATAC TTTTCTTAAAAATCA	MLLYCNPIYMKSSFPP KEIMRRNCTLVTEFILLGL TSRRELQILLFTLFLAIYM VTVAGNLGMIVLIQANA WLHMPMYFFLSHLSFVD LCFSSNVTPKMLEIFLSEK KSISYPACLVQCYLEIALV HVEIYLAVMAFDRYMAI CNPLLYGSRMSKSVCSFL ITVPYVYGALTGLMETM WTYNLAFCGPNEINHFC ADPLIKLACSDTYNKL SMFIVAGWNLSFSLFICIS YLYIFPAILKIRSTEGRQK AFSTCGSHLTAVTIFYAT LFFMYLRPPSKESVEQKG MVAVFYTTVIPMLNLIY SLRNKNVKEALIKELSMK IYFS

Table 1

Acc. No.	SEQ ID NO (Nuel) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
CG50267-01	15/16	CATGGGAAGATGGGTGAACCAAGTCTACACAGATGGCTTCTTCCCTCTTAGGCATC TTTTCCACAGCCAGACTGACCTTGCTCTTCTCTGCAGTTATGGTGGTCTTCAC AGTGGCCCTCTGTGGGAATGTCCCTCCTCATCTTCCCTCATCTACCTGGACGCTGGAC TTCACACCCCCATGTACTTCTTCCCTCAGCCAGCTCTCCCTCATGGACCTCATGTTG GTCTGTAAACATTGTGCCAAAGATGGCAGCCAACTTCCCTGTCTGGCAGGAAGTCCA TCTCCTTTGTGGCTGTGGCATAACAAATGGCTTTTGTGCTCTCTTGTGGGATCT GAGGGCTCTTGTCTGGACTCATGGCTTATGACCGCTACGTGGCCGTTAGCCACC CACTTCACTATCCCATCCTCATGAATCAGAGGGTCTGTCTCCAGATTACTGGGAG CTCTGGGCCCTTGGGATAATAGATGGAGTGATTCAGATGGTGGCAGCCATGGGC TTACCTTACTGTGGCTTGAGGAGCGTGGATCACTTTTCTGTGAGGTACAAAGCTT TATTGAAGCTGGCCGTGTCAGACACTTCCCTTTTGTGACACCCCTCTTGTGCTTGC TGTGCTTTCATGCTTCTCCTTCCCTTCTCCATCATCATGGCCTCCTATGCTTGCATC CTAGGGGCTGTGCTCCGAATACGCTCTGCTCAGGCCTGGAAAGCCCTTGGCCA CCTGCTCCTCCACCTAACAGCTGTACCCCTTCTATGGGCAGCCATGTTTCATG TACCTGAGGCCTAGGCGCTACCGGCCCCCTAGCCATGACAAGGTGGCCTCTATCT TCTACACAGTCTTACTCCCATGCTGAACCCCTCATTTACAGCTTGAGGAAATGG GGAGGTGATGGGGGCACTGAGGAAGGGGCTGGACCGCTGCAGGATTGGCAGCCA GCACTGAAC	MGRWVNQSYTDGFFLLG IFSHSQTDLVLFSAVMVV FTVALCGNVLLIFLIYLD GLHTPMYFFLSQLSLMDL MLVCNIVPKMAANFLSG RKSISFVGGCIQIFFVSL VGSEGLLLGLMAYDRYV AVSHPLHYPILMNQRVCL QITGSSWAFGIIDGVIQM VAAMGLPYCGLRSVDHF FCEVQALLKLACADTSLF DTLLFACCVFMLLLPFSII MASYACILGAVLRIRSAQ AWKKALATCSSHLTAVT LFYGAAMFMYLRPRRYR APSHDKVASIFYTVLTPM LNPLIYSLRNGEVMGALR KGLDRCRIGSQH

Table 1

Acc. No.	SEQ ID NO (Nuc) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
CG56818-01	17/18	GTTCCTGCAACTTCACACATGCCACCTTTGTGCTTATTGGTATCCCAGGATTAGAG AAAGCCCAATTCCTGGGTGGCTTCCCCCTCCTTCCATGTATGTAGTGGCAATGTT TGGAAACTGCATCGTGGTCTTCATCGTAAGGACGGAACGCAGCCTGCACGCTCCG ATGTACCTCTTTCTCTGCATGCTTGACGCCATTGACCTGGCCTTATCCACATCCAC CATGCCTAAGATCCTTGCCCTTTTCTGGTTGATTCCTCGAGAGATTAGCTTTGAGG CCTGCTTACCCAGATGTTCTTTATTTCATGCCCTCTCAGCCATTGAATCCACCATC CTGCTGGCCATGGCCTTTGACCGTTATGTGGCCATCTGCCACCCACTGCGCCCATG CTGCAGTGTCAACAAATACAGTAACAGCCCAAGATTGGCATCGTGGCTGTGGTCCG CGGATCCCCTCTTTTTCCTCCACTGCCTCTGCTGATCAAGCGGCTGGCCTTCTGCC ACTCCAATGTCTCTCGCACTCCTATTGTGTCCACCAAGGATGTAATGAAGTTGGC CTATGCAGACACTTTGCCCAATGTGGTATATGGTCTTACTGCCATTCTGCTGGTCA TGGCGGTGGACGTAATGTTCACTCTCTTGTCTTATTTCTGATAATACGAACGGTT CTGCAACTGCCCTCCAAGTCAGAGCGGCCCAAGGCCTTTGGAACTTGTGTGTAC ACATTGGTGTGGTACTCGCCTTCTATGTGCCACTTATTGGCCTCTCAGTTGTACAC CGCTTTGGAAACAGCCTTCATCCCATTTGTGCGTGTGTGATGGGTGACATCTACCT GCTGCTGCCCTCCTGTCAATCAATCCCATCACTATGTTGGTCCCAAAACCAACAGATC AGAACACGGGTGCTGGCTATGTTCAAGATCAGCTGTGACAAGGACTTGCAGGCT GTGGGAGGCAAGTGACCCTTAACACTACACTTCTCCTTATCTTTATTGGCTTGATA AACATAATTATTCT	SCNFTHATFVLGIPGLEK AHFWVGFPLLSMYVVA MFGNCIVVFIVRTERSLH APMYLFLCMLAAIDLALS TSTMPKILALFWFDSREIS FEACLTMFFIHLSAIES TILLAMAFDRYVAICHPL RHAAVLNNTVTAQIGIVA VVRGSLFFFPLPLLIKRLA FCHSNVLSHSYCVHQDV MKLAYADTLPNVVYGLT AILLVMGVDVMFISLSYF LIIRTVLQLPSKSERAKAF GTCVSHIGVVVLAIFYVPLI GLSVVHRFGNSLHPVVRV VMGDIYLLLPVINPIIYG AKTKQIRTRVLAMFKISC DKDLQAVGGK

Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
CG57290-01	19/20	AATGGCTGCAGGAAATCACTCTACAGTGACAGAGTTCAATCTCAAAGGGTTAAACG AAGAGAGCAGACCTCCAGCTCCCCCTCTTTCTCCTCTTCTCGGGATCTACTTGGT CACCATCGTGGGAAACCTGGGCAATGATCACTCTAAATTTGTCTGAACTCTCAGCTG CACACCCCATGTACTACTTTCTCAGCAAATCTGTCACTCATGGACCTCTGCTACTC CTCCGTCATTACCCCTAAGATGCTGGTGAACTTTGTGTGTCAGAGAAAAACATCATC TCCTACGCAGGGTGCAATGTCACAGCTCTACTTCTTCTCCTTGTGTTTGTCAATTGCTGA GTGCTACATGCTGAGAGTGATGGCCTACGACCGCTATGTTGCCATCTGCCACCCCT TTGCTTTACAACATCATATTATGTCTCATCACACCTGCCTGCTGCTGGTGGCTGTGGT CTACGCCATCGGACTCATTTGGCTCCACAATAGAAACTGGCCTCATGTTAAAACTG CCCTATTGTGAGCACCTCATCAGTCACACTACTTCTGTGACATCCCTCCCTCATGAA GCTGCTCTCTAGCACCTATGATGTTGAGATGACAGTCTTCTTTTGGCTGGAT TCAACATCATAGTCACGAGCTTAACAGTCTTGTGTTCTTACACCTTCATTCTCTCC AGCATCCTCGGCATCAGCACCCACAGAGGGGAGATCCAAAGCCTTCAGCACCTGC AGTCCCACTTGCAGCCGTGGGAATGTTCTATGGATCAACTGCATTCATGTACT TAAACCCCTCCACAAATCAGTTCCTTGACCCAGGAGAAATGTGGCCTCTGTGTTCCA CACCACGGTAATCCCCCATGTTGAATCCCCTAATCTACAGCCTGAGGAACAAGGAA GTAAAGGCTGCCGTGCAGAAAAACGCTGAGGGGTAAACTGTTTGTGATGCAAA	MAAGNHSTVTEFILKGLT KRADLQLPLFLFLGIYL VTIVGNLGMITLICLNSQL HTPMYYFLSNLSLMDLC YSSVITPKMLVNFVSEKN IISYAGCMSQLYFFLVFVI AECYMLRVMMAYDRYVAI CHPLLNYNIIMSHHTCLLL VAVVYAIGLIGSTIETGL MLKLPYCEHLISHYFCDI LPLMKLSCSSTYDVEMT VFFLAGFNIIVTSLTVLVS YTFILSSILGISTTEGRSKA FSTCSSHLAAVGMFYGST AFMYLKPSTISSLTQENV ASVFHTTVIPMLNPLIYSL RNKEVKAAVQKTLRGKL F

Table 1

Acc. No.	SEQ ID NO (Nucl) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAL365336_G	21/22	TGATCATAAATGAATGTGAGCTTCAAGACTGGATTCTCTCTCATGGGGTTCTCTGA TGAGCGTAACCTTCAGATTTACATGCAGTGTCTCTTTTGATCACATACCTGTTGG CCATCATGGGCAATCTGCTCAATTATCACCATCATCACCTTGGACCAACGCTCTGCAT TCTCCCATGTACTACTTCTTGAAGCACCTCTCTTTTGGATCTCTGCTTCATCTCT GTTACTGTTCTCAGTCTATTGCAAACTCACTCATGAACAAATGGTTTCATTTCTCT TGGTCAGTGTATGCTTCAGGTTTTCTTCTTCATAGCTCTGGCTCATCAGAAGTAG CTATTCTCACAGTGTCTTATGACCGGTATGTTGCCATCTGTGGGCCACTGCAG TATGAGACAAATATGGATCCCCATGCCTGCAAGTGCACAGTGTAAATTTCTCAAATTCCTCTT TGGCTGGAGGACTATCTGGGCTCTACACACAGGTGTTAAATTTCTCAAATTCCTCTT TGTGGGAAGAGAAATTAATCACCAAGTTCTTCTGTGACATTCGCCCAAATGCTAAAC TAGCTTGTCTTATGAATTCATTAATGAGATTGACGTGGCTGCATTTACAAACATCC ACAGCCTTTGTCTGTTTAATAGCCATAGTCTTCTCTCTATACTCAGATCTTCTCAAC TGTGATGAGAAATCCATCAGCTGATAGTCGGACTAAAGGTGTTCTCCACCTGTCTA CCACATTTGTTGTAGTCAATGTTCTTCTCTCAGCTGCAGGCTTTGAAATTTCTAAG ACCTCCTTCAGATTCCCTGTGAGCAATGGACCTCGTATTCCTCATATTCTACACTG TGATACCTCCAAACACTCAAATCCACTCATCTACAGCTTGAGGAATGAGGCCATGAA AGCAGCTCTGAGGAAAGTGTGTCAAAGAAAGAAATTTTCTCGGAGAAATGGTATAT GTAAAGCTATATTCAATCTCTAAAGAGACAAACAACTAAGAGGCATTGCTACTA T	MNVSFKGTGFLLMGFSD RNLQILHAVLFLITYLLAI MGNLLIITIITLDQRLHSP MYYFLKHLSDLCLCFISV TVPQSIANSMLMNGFISL GQCMLQVFFFIALASSEV AILTVMSYDRYVAICRPL QYETIMDPHACKCAVIA WMAGLSGLLHTGVNFS IPLCGKRIIHQFFCDIPQM LKLACSYEFINEIAVAAFT TSTAFVCLIAIVFSYTQIFS TVMRIPSADSRTKVFSTC LPHLFVVMFFLSAAGFEF LRPPSDLSAMDVLVFSIFY TVIPPTLNPLIYSLRNEAM KAALRKVLSKEEFSRRM VYVKAIFNL









Table 1

Acc. No.	SEQ ID NO (NucI) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAL359352_D	29/30	ATGAGTATCAAACTGCTCTCTGTGGCAGGAGAACAGCTTGCTGTCAACACGCTTTG CATTITCCAAAGTTCTCTGAGGTCCCTGGAGAAATGCTTCTCTCTCTGTTACCCATCATC CTCCTCATGTTCTTAGTATCTCTGACAGGAAATGAACATCATAGCCATTGCCATCTG CACCAGTCCAGCCCTACATACCCCCATGTACTTCTTCTAGCCCAACTGTCTCTTC TGGAGATTGGTACACTTGTCTGTCTATACCCAAATGCTACAGAGCCTTGTAAG TGAGGCCCGAGAAATCTCTCGGGAGGGATGTGCCACACAGATGTTTCTTTCACA TTTTITGGTATAACTGAGTGTCTACTGGCAGCCATGGCCTATGACCGCTGCA TGGCCATATGCTCCCCACTCCACTATGCAACACGAATGAGTCATGGGGTATGTGC CCATTGGCAATAGTTTCATGGGGAATGGGATGTATAGTAGGGTTGGACAAACC AATTTTATTTCTCGTTGAACTTCTGTGGACCTGTGAGATAGACCACTTCTTCTG TGACCTTCCACCTGTCTGGCACTTGCCTGTGGAGATACTTCCCAAAATGAGGCT GCAATCTTCGTGGCAGCAATCTTTCATATCTAGCCCCATTTTGTGTGATCATTTA TTCCTATGTCAGAAATCTGGTTCAGTGTCTGATGCTTACCTTCACCTGAGGGCGC CATAAAGCTCTCTCCACCTGTTCCTCGCATCTACTGTAGTCACACTGTTTTTGG CTCAGGATCTATTACCTACTTGAGGCCCAAGTCTAGCCACTTACCAAGGAATGGAC AAACCTCTGGCCCTTTCTACACCGCGGTGACATCCATGCTGAACCCCATCATCTA TAGCTTAAGGAACAAGGAAGTGAAGACAGCACTGAGAAAACACTGAGTCTGAA GACATCTCGGGCAATAAAATAGGTAACAGAACCTTGCAGAGCTGCTGGCTAATGA AAAT	MSINCSLWQENSLSVKRF AFSKFSEVPGECLFLTLI LLMFLVSLTGNELIAIAC TSPALHTPMYFFLANLSL LEIGYTCVPIKMLQSLVS EAREISREGCATQMFFFT FFGITECCLLAAMAYDRC MAICSPLHYATRMISHV CAHLAIVSWGMCIVGL GQTNFIFSLNFCGPCEIDH FFCDLPPVLALACGDTSQ NEAAIFVAAILCISPFLLII YSYVRILVAVLVMPSPG RHKALSTCSSHLLVVTLF FGSGSITYLRPKSSHLPG MDKLLALFYTAVTSMNL PIYSLRNKEVKTALRKTL SLKTSRAINR

Table 1

Acc. No.	SEQ ID NO (Nuc) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAC036111_C_31/32		GAAACATGTTTCTGACAGAGAGAAAATACGACATCTGAGGCCACATTCACTCTCT TGGGCTTCTCAGATTACCTGGAACTGCAAAATCCCTCTTCTTTGTATTTCTGGCA GTCTACGGCTTCAGTGTGGTAGGAATCTTGGGATGATAGTGATCATCAAAATTA ACCCAAAATTGCATACCCCATGTATTTTCTCAACCACTCTCCTTTGTGGAT TTCTGCTATTCTCTCCATCATGCTCCCATGTATGCTGTGAACCTGGTTGTAGAAGA TAGAACCAATTCATCTCAGGATGTTGGTGCAATCTTTTCTTTTGCACCTTTG TAGTGAATGAATTAATCTATTGCGGTGATGGCTATGACCACTTTGTGGCCATT TGCAATCCTCTGCTACACAGTTGCCATCTCCAGAAACTCTGTGCCATGCTGCT GGTTGATTGTATGCATGGGGAGTGGCATGTTCCCTGACACTCGCGTCTCTGCT TTAAAGTTATCTTTTCAATGTTCAACACAAATCAATCAATTTCTTCTGTGAGTTATC CTCCCTGATATCACTCTTACCTGACTCTTATCTCAGCCAGTTGCTTCTTTTCA TGTTGCCACTTTTAAATGAGATAAGCACACTACTCATCTTCTGACATCTTATGCAT TCATCATTTGTACCACTTGAAGATGCCTTCAGCCAGTGGGCAACCGCAAGTCTT CTCCACCTGTGCTCCCATGACTGCCATCACCATCTTCCATGGCACCATCTCTCT TCCTCTACTGTGTAACCACTCCAAAACCTCCAGGCACACAGTCAAAAGTGGCCTC TGTGTTTACACCGTGGTGATCCCTTGTGAAATCCCTGATCTACAGTCTGAGAA ATAAAGATGTTAAGGATGCAATCCGAAAAATAATCAATACAAATATTTTCATAT TAAACATAGGCATTGGTATCCATTAAATTTTGTATTGAACAAATAAATTTTCTCT	MFLTERNTTSEATFTLLG FSDYLELQIPLFFFLAVY GFSVVVGNLGMIVIKINPK LHTPMYFFLNHLFSVDFC YSSIIAPMMLVNLVVEDR TISFSGCLVQFFFEFTFVV TELILFAVMAVDHFVAIC NPLLVTVAISQKLCAMLV VVLVYAWGVACSLTLACS ALKLSFHGFNTINHHFCEL SSLISLSPDSYLSQLLFT VATFNEISTLLIILTSYAFII VTTLKMPSASGHRKVFST CASHLTATIFHGTILFLY CVPNSKNSRHTVKVASV FYTVVIPLLNPLIYSLRNK DVKDAIRKIINTKYFHIKH RHWYPFNFVIEQ

Table 1

Acc. No.	SEQ ID NO (Nuc) / SEQ ID NO (Prot)	DNA SEQUENCE	PROTEIN SEQUENCE
GMAC069559_F	33/34	AATGGACTCAGTAAATGTCTCTCTGGTGACTGAATTCCTTCTAGTAGGATTAACA CATCAGCCTGATCGCCAAATACCCCTGTCTCTCTGTTTCTAGCAATGTATCTAGT CACTGCAATTGGGAAATTTGGGTTTGATTATCTAGTTTGTCTAAATTCACATCTTC ACACACCATGTACTTTTTCTCTTTAACTTGTCCTTTATAGATTTTGTGTTATTCTT CTGTGTTCACTCCAAAAATGTTGATGAACCTTCATATTAAAGACAGAAATGCCATTTT CTATATGCAATGATGACTCAGCTCTACTCTTTTGTGTTTGTGTTGTTCTGAAT GCTTTGTGCTGACGTCAATGGCTTATGATCGATATGTGGCTATCTGTAATCCACTT TTATATAATGTGATGATTTCCCTCAAGTGTGTTAAACCTAATGATTGGTTCTTA TTTGATGGCAATTTCTGAGGCTGTAGCTCTCACTGTGTGATGCTGACATTGACTT TCTGTGATGGAACATCAACCACTACTTCTGTGACATCCTTGCTCTGTTCCAGCTC TCCTGCTCAAGCACCTATGTTAATAAGCTTGTAGCTTATGTCTATGCTCTCCAGCA ACATACTTTTCTACTCTACCATCTTTATCTCTTATGGTTTATCCTCTCCAGCA TCTTCCGCATAAGTTCCTCTAAGGGTAGGTCCAAAGCCTTCAGCACCTGCAGCTC CCACATAATCGCTGTTCTCTGTTCTTTGGTTCAGGAGCATTTGTGATTTCAAAC CCTCCTCACCTGGGTCTATGGAATGGCAAAAATCTCTCTGTTTATACCAAT GTAGTTCCCTATGATGAAATCCATTAATCTACAGCTTGAAAAACAAGATGTTAAAA TTGCCCTAAGAAAAATCTTTGGCAAGGTGGAAGATTGATTGGATACATAT	MDSVNVSLVTEFLLVGLT HQPDRIPLFLFLAMYL VTALGNLGLIILVLLNSHL HTPMYFFLFNLSFIDFCYS SVFTPKMLMNFILRQNAI SYMQCMTQLYFFCFV SECFVLTSMAYDRYVAIC NPLLNVVMISPPQVCLNL MIGSYLMAFSEAVALTVC MLTLTFCDGNINHYFCDI LALFQLSCSSTYVKNLVA YVIVVINILFSTPTIFISYG FILSSIFRISSSKGRSKAFS TCSSHIAVSLFFGSGAFV YFKPSSPGSMEWAKISSV FYTNVVPMMNPLIYSLK NKDVKIALRKSLARWKI

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## OTHER EMBODIMENTS

Although particular embodiments have been disclosed herein in detail, this has been done by way of example for purposes of illustration only, and is not intended to be limiting with respect to the scope of the appended claims, which follow. In particular, it is  
5 contemplated by the inventors that various substitutions, alterations, and modifications may be made to the invention without departing from the spirit and scope of the invention as defined by the claims. The choice of nucleic acid starting material, clone of interest, or library type is believed to be a matter of routine for a person of ordinary skill in the art with knowledge of the embodiments described herein. Other aspects, advantages, and  
10 modifications considered to be within the scope of the following claims.